=> FILE REG

L20

1010 SEA LAVAL#

FILE 'REGISTRY' ENTERED AT 14:30:25 ON 20 DEC 2006
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## => DISPLAY HISTORY FULL L1-

-> Didimii iiididki idhi hi		
	FILE 'REGI	STRY' ENTERED AT 11:46:53 ON 20 DEC 2006 E HYDROGEN/CN
L1	1	SEA HYDROGEN/CN
		E OXYGEN/CN
L2	1	SEA OXYGEN/CN
	•	E WATER/CN
L3	1	SEA WATER/CN
		E IODINE/CN
L4	1	SEA IODINE/CN
		E NITROGEN/CN
L5	1	SEA NITROGEN/CN
		E PHOSPHORIC ACID/CN
L6	1	SEA "PHOSPHORIC ACID"/CN
		LUS' ENTERED AT 11:50:55 ON 20 DEC 2006
L7	5050	SEA BAR ?/AU OR BARGADDA ?/AU OR BAR GADDA ?/AU OR GADDA
т о	001007	?/AU OR GADDA BAR ?/AU
L8		SEA HYDROGEN#/TI
L9 L10		SEA PLASMA#/IT SEA L7 AND L8 AND L9
L11		SEA L7 AND L6 AND L9 SEA L3 (L) RACT/RL
L12		SEA L10 AND L11
птг	U	SEA DIO AND DII
	FILE 'HCA'	ENTERED AT 11:56:37 ON 20 DEC 2006
L13		SEA L1/P
L14		SEA L2/P
L15		SEA L3 OR STEAM? OR (WATER? OR H2O OR HOH) (2A) (VAPOUR?
		OR VAPOR? OR EVAP? OR VOLATILIZ? OR VOLATILIS? OR
		ATOMIS? OR ATOMIZ? OR MIST OR MISTS OR MISTED OR
•		MISTING# OR SPRAY? OR MICRONIZ? OR MICRONIS? OR GAS## OR
		GASEOUS? OR GASIF?)
L16	130639	SEA L4 OR I2 OR (MOLECULAR? OR ELEMENTAL? OR PURE# OR
		PURIF?) (2A) (IODINE# OR I)
L17	462901	SEA L5 OR N2 OR (NITROGEN# OR N) (2A) (GAS## OR GASEOUS?
		OR GASIF? OR ATM# OR ATMOS?)
L18	147311	SEA L6 OR (PHOSPHORIC# OR ORTHOPHOSPHORIC#)(2A)ACID# OR
		H3PO4
L19		QUE MEMBRAN?
		A

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L21
         107425 SEA MICROWAV?
L22
         107338 SEA RADIOFREQ? OR RADIO#(2A)(FREQ# OR FREQUENC?) OR RF
                OR R(W)F
          55749 SEA (ELECTRIC? OR ELEC# OR DISCHARG?) (2A) (ARC OR ARCS)
L23
                OR (GLOW? OR SPARK?) (2A) DISCHARG?
          37275 SEA (ELECTROMAG? OR ELECTRO(A) (MAGNET? OR MAG#) OR EM OR
L24
                E(W)M)(2A)(FIELD? OR FLUX? OR WAVE# OR WAVING#)
L25
                QUE PLASM#
L26
          32996 SEA L3 (L) (PROC# OR PROCESS?)
L27
           4036 SEA L13 AND L14
L28
            382 SEA L27 AND L26
L29
              8 SEA L28 AND L25
L30
           2305 SEA L27 AND L15
L31
             43 SEA L30 AND L25
              0 SEA L31 AND L16
L32
L33
              5 SEA L31 AND L17
L34
              2 SEA L31 AND L18
              5 SEA L31 AND L19
L35
L36
             0 SEA L31 AND L20
L37
             6 SEA L31 AND ((L21 OR L22 OR L23 OR L24))
L38
             13 SEA (L32 OR L33 OR L34 OR L35 OR L36 OR L37)
         319842 SEA L1
L39
L40
         394274 SEA L2
          40452 SEA L39 AND L40
L41
L42
           9809 SEA L41 AND L15
L43
            532 SEA L42 AND L25
L44
            316 SEA L43 AND ((L16 OR L17 OR L18 OR L19 OR L20 OR L21 OR
                L22 OR L23 OR L24))
L45
              8 SEA L43 AND L16
L46
            241 SEA L43 AND L17
L47
             5 SEA L43 AND L18
L48
            24 SEA L43 AND L19
L49
              0 SEA L43 AND L20
L50
            110 SEA L43 AND ((L21 OR L22 OR L23 OR L24))
L51
             40 SEA L43 AND L21
L52
             30 SEA L43 AND L22
L53
             49 SEA L43 AND L23
              8 SEA L43 AND L24
L54
L55
             4 SEA L48 AND (L51 OR L52 OR L53)
L56
            23 SEA L50 AND L26
L57
             3 SEA L46 AND L47
             18 SEA L29 OR L38
L58
L59
            25 SEA L31 NOT L58
L60
              0 SEA L59 AND (L45 OR L47 OR L48 OR L54 OR L55 OR L56 OR
                L57)
             21 SEA (L45 OR L47 OR L54 OR L55 OR L57) NOT (L58 OR L59)
L61
            36 SEA (L48 OR L56) NOT (L58 OR L59 OR L61)
L62
L63
            16 SEA 1840-2003/PY, PRY AND L58
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L64 22 SEA 1840-2003/PY,PRY AND L59 L65 19 SEA 1840-2003/PY,PRY AND L61 L66 32 SEA 1840-2003/PY,PRY AND L62

## => FILE HCA

FILE 'HCA' ENTERED AT 14:30:44 ON 20 DEC 2006
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## => D L63 1-16 CBIB ABS HITSTR HITIND

L63 ANSWER 1 OF 16 HCA COPYRIGHT 2006 ACS on STN

142:264323 Ecology system focusing on hydrogen use. Yoshino, Kazutora (USA). U.S. Pat. Appl. Publ. US 2005044853 A1 20050303, 14 pp. (English). CODEN: USXXCO. APPLICATION: US 2003-653553 20030902.

AB Ecosystem that is highly efficient and environmental is shown. Turbo Hydrogen engine with controlled combustion using the recycling of uncombusted gases such as hydrogen and oxygen can be embedded to this ecosystem. Solar energy and wind energy can be used as external energy source. The energy of motion of a vehicle is charged back to usable energy when it reduces the motion.

IT 7732-18-5, Water, processes

(ecol. system focusing on hydrogen use)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

o = o

IT 1333-74-0P, Hydrogen, uses (ecol. system focusing on hydrogen use)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

```
IC
     ICM F03G006-00
INCL 060641800
     52-1 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     Section cross-reference(s): 59, 71, 72
IT
     Fission
     Nuclear fusion
       Plasma
        (engine; ecol. system focusing on hydrogen use)
IT
        (phosphoric acid; ecol. system focusing on
        hydrogen use)
     Fuel cells
IT
        (proton exchange membrane; ecol. system focusing on
        hydrogen use)
IT
     64-17-5, Ethanol, processes
                                   67-56-1, Methanol, processes
     142-82-5, Heptane, processes 7732-18-5, Water,
        (ecol. system focusing on hydrogen use)
IT
     7782-39-0P, Deuterium, preparation 7782-44-7P, Oxygen,
                   10028-17-8P, Tritium, preparation
     preparation
                                                      12586-59-3P,
     Proton
        (ecol. system focusing on hydrogen use)
IT
     1333-74-0P, Hydrogen, uses
        (ecol. system focusing on hydrogen use)
     ANSWER 2 OF 16 HCA COPYRIGHT 2006 ACS on STN
142:117686 Integrated plasma fuel cell process.
                                                   Steinberg,
     Meyer (USA). PCT Int. Appl. WO 2005004255 A2 20050113, 16 pp.
     DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR,
     BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG,
     ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP,
     KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
     NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,
     SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW;
     RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA,
     GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR.
     (English).
                 CODEN: PIXXD2. APPLICATION: WO 2004-US20019 20040624.
     PRIORITY: US 2003-2003/604153 20030627; US 2003-2003/604961
     20030828.
AB
     The invention concerns a method for efficiently producing energy,
     carbon, carbon monoxide, synthetic carbonaceous liq. and gaseous
     fuels and hydrogen from fossil or biomass fuels with minimal carbon
     dioxide and other emissions.
IT
     7732-18-5, Water, processes
        (integrated plasma fuel cell process)
RN
     7732-18-5 HCA
```

Water (8CI, 9CI) (CA INDEX NAME)

CN

```
H<sub>2</sub>O
IT
     1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen,
     uses
         (integrated plasma fuel cell process)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI)
CN
                         (CA INDEX NAME)
o = 0
     ICM H01M
IC
CC
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     Section cross-reference(s): 51, 72
ST
     plasma fuel cell process
IT
     Hydrocarbons, uses
         (C1-4; integrated plasma fuel cell process)
     Thermodynamic cycle
IT
         (Rankine; integrated plasma fuel cell process)
IT
     Fuel cells
        (direct carbon; integrated plasma fuel cell process)
IT
     Fuels
        (fossil; integrated plasma fuel cell process)
     Carbonaceous materials (technological products)
IT
        (fuels; integrated plasma fuel cell process)
     Biomass
IT
     Diesel fuel
     Electrolytic cells
     Fischer-Tropsch reaction
       Plasma
     Water gas shift reaction
        (integrated plasma fuel cell process)
IT
     Gasoline
        (integrated plasma fuel cell process)
IT
     Fuel cells
        (solid oxide; integrated plasma fuel cell process)
IT
     Boilers
        (steam; integrated plasma fuel cell process)
     7732-18-5, Water, processes
       (integrated plasma fuel cell process)
```

```
LANGEL 10/632,708
IT
     67-56-1, Methanol, uses
                               74-82-8, Methane, uses
        (integrated plasma fuel cell process)
IT
     124-38-9P, Carbon dioxide, uses 630-08-0P, Carbon monoxide, uses
     1333-74-0P, Hydrogen, uses
                                  7440-44-0P, Carbon, uses
     7782-44-7P, Oxygen, uses
        (integrated plasma fuel cell process)
    ANSWER 3 OF 16 HCA COPYRIGHT 2006 ACS on STN
142:117577 Hydrogen generation by water splitting in a radiant energy
     transfer reactor. Bar-Gadda, Ronny (Bar-Gadda, LLC, USA).
     Appl. WO 2005005009 A2 20050120, 35 pp. DESIGNATED STATES: W:
     AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO,
     CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM,
     HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
     LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH,
     PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ,
     UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG,
    CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML,
    MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2.
    APPLICATION: WO 2004-US21267 20040630. PRIORITY: US
     2003-2003/PV48408U 20030630; US 2003-2003/632708 20030801; US
     2004-2004/819591 20040406.
    Hydrogen is produced by introducing high temp. steam or
AΒ
    water vapor into a radiant energy transfer
              The radiant energy is absorbed by the mols. which dissoc.
     into their constituent mol. elements of hydrogen and oxygen.
    prevent recombining of the constituent mol. elements, the hydrogen
     and oxygen are sepd. from each other by a membrane,
     centrifugation using a variant electromagnetic
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field, or by chem. reaction of one component. The source of steam can be geothermal steam, or steam produced in a boiler using heat from combustion processes. Once sepd., the mol. components are prevented from recombining with each other or with other elements by using std. sepn. techniques normally employed for sepg. dissimilar gaseous species.

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation

(hydrogen generation by water splitting in radiant energy transfer reactor)

RN1333-74-0 HCA

Hydrogen (8CI, 9CI) (CA INDEX NAME) CN

H-H

RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) o = 0IT 7732-18-5, Water, processes (hydrogen generation by water splitting in radiant energy transfer reactor) RN7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME) H20 IT 7664-38-2, Phosphoric acid, reactions 7727-37-9, Nitrogen, reactions (hydrogen generation by water splitting in radiant energy transfer reactor) 7664-38-2 HCA RN CN Phosphoric acid (7CI, 8CI, 9CI) (CA INDEX NAME) HO- P-OH OH 7727-37-9 HCA RN CNNitrogen (8CI, 9CI) (CA INDEX NAME) ICM B01D IC CC 52-1 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49 hydrogen manuf water splitting plasma STIT Combustion enthalpy Electric arc Electromagnetic field Heat transfer Plasma Turbines Waste plastics and rubbers (hydrogen generation by water splitting in radiant energy

transfer reactor)

IT

1333-74-0P, Hydrogen, preparation 7782-44-7P,

Oxygen, preparation

(hydrogen generation by water splitting in radiant energy transfer reactor)

IT 7732-18-5, Water, processes

(hydrogen generation by water splitting in radiant energy transfer reactor)

IT 124-38-9, Carbon dioxide, reactions 7664-38-2,

Phosphoric acid, reactions 7727-37-9,

Nitrogen, reactions

(hydrogen generation by water splitting in radiant energy transfer reactor)

L63 ANSWER 4 OF 16 HCA COPYRIGHT 2006 ACS on STN

141:280318 Water dissociator for hydrogen use in engines. Konnov, S. V. (Russia). Russ. RU 2230916 C2 20040620, No pp. given (Russian). CODEN: RUXXE7. APPLICATION: RU 2002-116938 20020624.

AB A proposed water dissociator makes it possible to dissoc. water periodically into hydrogen-oxygen gas mixt., expecially for use in engines. The water dissociator contains a nozzle installed in an internal combustion engine cylinder and connected through a pump to a water tank. A device for coupling with a superhigh frequency electromagnetic oscillator is installed in the engine cylinder. A high-speed two-stroke carburetor internal combustion engine is used. This method improves the possibility of dissocn. of water for use in engines.

IT 7732-18-5, Water, uses

(water dissociator for hydrogen use in engines)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H20

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation

(water dissociator for hydrogen use in engines)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0==0

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IC ICM F02B047-02
ICS F02M025-03
```

- CC 52-1 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49
- ST water dissociator **microwave** oscillator internal combustion engine **plasma** fuel
- IT Cold plasma

(created inside engine cylinder; water dissociator for hydrogen use in engines)

IT Internal combustion engines

Microwave oscillators

Pistons

Valves

(water dissociator for hydrogen use in engines)

IT 7732-18-5, Water, uses

(water dissociator for hydrogen use in engines)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P,
Oxygen, preparation
(water dissociator for hydrogen use in engines)

L63 ANSWER 5 OF 16 HCA COPYRIGHT 2006 ACS on STN

- 140:409609 Method for gasification of carbon-containing substances by using plasma. Stari, Johannes; Simmel, Johannes (Johann Stari Ges. m.b.H. Sonder- und Spezialmaschinenbau, Austria). PCT Int. Appl. WO 2004041974 A1 20040521, 25 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (German). CODEN: PIXXD2. APPLICATION: WO 2003-AT315 20031020. PRIORITY: AT 2002-1662 20021104.
- The invention relates to a method for qasification of carbon-contq. AΒ substances. The substances are fed into the reactor through a transfer channel. Plasma torches for generating a gaseous plasma are arranged at the reactor bottom in the direction of gravitation force. The oxidn. products, i.e. clinker, glass, ashes are drawn downward and fed into a solidifier, inside of which the oxidn. products are sepd., subsequently collected in the collecting tank, and transported away when required. The gas produced inside the reactor is fed via line into the gas washer, inside of which the gas is cooled and purified with sprayed water. The discharging water is charged via line into the collecting tank, inside of which water is already present so that the clinker, glass or the like can be cooled. The purified gas is fed via line into the compressor, inside of which the gas is

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compressed to 2 bar and then purified once more while passing through filter, and is then mixed inside the gas mixer with atm. This mixt. of gas and air is then fed via line into the gas engine that drives a generator. 1333-74-0P, Hydrogen, preparation 7727-37-9P, Nitrogen, preparation 7782-44-7P, Oxygen, preparation (method for gasification of carbon-contg. substances by using plasma) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7727-37-9 HCA Nitrogen (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0ICM C10J003-46 ICS C10J003-52; B01J019-08; C10J003-18 52-1 (Electrochemical, Radiational, and Thermal Energy Technology) fuel gas manufg wood chip gasification plasma Abies Fagus Larix Wood (chips; method for gasification of carbon-contg. substances by using plasma) Plasma Plasma torches (for gasification of carbon-contg. substances by using plasma) Fuel gas manufacturing (gasification, fluidized-bed; method for gasification of carbon-contg. substances by using plasma) Fuel gas manufacturing (gasification; method for gasification of carbon-contg. substances by using plasma)

IT 124-38-9P, Carbon dioxide, preparation 630-08-0P, Carbon monoxide,
 preparation 1333-74-0P, Hydrogen, preparation
 7727-37-9P, Nitrogen, preparation 7782-44-7P,
 Oxygen, preparation
 (method for gasification of carbon-contg. substances by using
 plasma)

ANSWER 6 OF 16 HCA COPYRIGHT 2006 ACS on STN L63 Electrochemical generation, storage and reaction of hydrogen 140:273630 Sanders, Nicholas (Diffusion Science, Inc., USA). Int. Appl. WO 2004027901 A2 20040401, 92 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2003-US29802 20030917. PRIORITY: US 2002-411443P 20020917; US 2003-455215P 20030317.

The invention concerns an electrolytic app. for using AB . catalyst-coated hollow microspheres to produce gases, store them, and to make them available for later use. The app. uses catalyst-coated hollow microspheres in reversible electrochem. processes and reactions, such as those used in conjunction with water dissocn., fuel cells, and rechargeable batteries. can be used to manuf. and store hydrogen and or oxygen and to make them available for subsequent use as raw materials for use in electrochem. and chem. reactions or as a fuel and or oxidizer for a combustion engine. The app. can be used as a hydrogen-oxygen hermetically sealed secondary battery. The app. can be used as a hydrogen storage portion of certain types of secondary batteries. Hydrogen and oxygen can be stored within hollow microspheres at moderate temp. and pressure, eliminating the need for expensive storage and handling equipment, and increasing the mobility of hydrogen-powered vehicles. Storage of hydrogen and or oxygen within the microspheres significantly reduces flammability and explosion concerns and resolves many fuel cell scalability issues.

IT 7732-18-5, Water, processes

(electrochem. generation, storage and reaction of hydrogen and oxygen)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P,

Oxygen, preparation

(electrochem. generation, storage and reaction of hydrogen and oxygen)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IC ICM H01M004-00

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 57, 72

IT Vapor deposition process

(plasma; electrochem. generation, storage and reaction of hydrogen and oxygen)

IT 7732-18-5, Water, processes

(electrochem. generation, storage and reaction of hydrogen and oxygen)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P,

Oxygen, preparation

(electrochem. generation, storage and reaction of hydrogen and oxygen)

- L63 ANSWER 7 OF 16 HCA COPYRIGHT 2006 ACS on STN
- 139:166829 Hydrogen generation from water, methane, and methanol with nonthermal plasma. Kabashima, Hajime; Einaga, Hisahiro; Futamura, Shigeru (National Institute of Advanced Industrial Science and Technology, Ibaraki, 305-8569, Japan). IEEE Transactions on Industry Applications, 39(2), 340-345 (English) 2003. CODEN: ITIACR. ISSN: 0093-9994. Publisher: Institute of Electrical and Electronics Engineers.
- AB Hydrogen generation from water, methane, and methanol was studied with different types of nonthermal plasma reactors under different conditions. With a ferroelec. packed-bed reactor in N2, hydrogen gas yield decreased in the order: methanol > methane > water. A similar trend was obsd. with a silent discharge plasma reactor, but H2 yields were much lower with the latter reactor. At fixed specific energy densities, higher H2 yields were obtained at higher gas flow rates in the reactions of the above substrates. The initial water concn. was optimized at .apprx.2.0% to obtain the highest rate for H2 formation. Under the

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H<sub>2</sub>O

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same conditions, H2 yield decreased in the order: Ar > N2 > air ≈ O2. The ferroelec. packed-bed reactor could be operated continuously for 10 h without any decrease in its performance in the H2 generation from water. 1333-74-0P, Hydrogen, preparation (hydrogen generation from water, methane, and methanol with nonthermal plasma) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) H-H7782-44-7P, Oxygen, preparation (hydrogen generation from water, methane, and methanol with nonthermal plasma) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0**7727-37-9**, Nitrogen, uses (hydrogen generation from water, methane, and methanol with nonthermal plasma) 7727-37-9 HCA Nitrogen (8CI, 9CI) (CA INDEX NAME) **7732-18-5**, Water, reactions (hydrogen generation from water, methane, and methanol with nonthermal plasma) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 52-1 (Electrochemical, Radiational, and Thermal Energy Technology) hydrogen water methane methanol nonthermal plasma ferroelec bed packing; plasma silent discharge reactor hydrogen controlled atm effect

(C2; hydrogen generation from water, methane, and methanol with

Hydrocarbons, formation (nonpreparative)

nonthermal plasma)

IT Air

Controlled atmospheres

Decomposition

Ferroelectric materials

Plasma

Reaction mechanism

(hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT Reactors

(packed-bed; hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT Reactors

(plasma; hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT Electric discharge

(silent, tubular; hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT 12047-27-7, Barium titanium trioxide, uses (bed packing; hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT 630-08-0P, Carbon monoxide, preparation (hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT 67-56-1, Methanol, processes (hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT 74-84-0, Ethane, formation (nonpreparative) 74-85-1, Ethene, formation (nonpreparative) 74-86-2, Acetylene, formation (nonpreparative) 124-38-9, Carbon dioxide, formation (nonpreparative)

(hydrogen generation from water, methane, and methanol with nonthermal plasma)

IT 7440-37-1, Argon, uses 7727-37-9, Nitrogen, uses (hydrogen generation from water, methane, and methanol with nonthermal plasma)

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L63 ANSWER 8 OF 16 HCA COPYRIGHT 2006 ACS on STN
131:292518 Electricity feeders for water electrolytic cells. Shinohara,
     Taizo; Yamaguchi, Mikimasa (Fuji Electric Corporate Research and
     Development, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 11302891 A2
     19991102 Heisei, 4 pp. (Japanese).
                                          CODEN: JKXXAF.
     APPLICATION: JP 1998-114476 19980424.
AB
     The electricity feeder is made of a Ti fiber-sintered plate whose
     surface is made smooth by plasma thermal spray of Ti in an
     inert gas (Ar, He, or N) atm. The durability of
     membrane electrode joined body is improved thus efficient
     water electrolytic cells can be provided.
IT
    7727-37-9, Nitrogen, uses
        (electricity feeder made of a Ti fiber-sintered plate whose
        surface is made smooth by plasma thermal spray of Ti in
        an inert gas (Ar, He, or N) atm.)
RN
     7727-37-9 HCA
CN
     Nitrogen (8CI, 9CI) (CA INDEX NAME)
IT
     7732-18-5, Water, processes
        (electricity feeders for water electrolytic cells)
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME).
CN
H<sub>2</sub>O
IT
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
     Oxygen, preparation
        (electricity feeders for water electrolytic cells for prodn. of)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
o = o
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IC

ICM C25B009-04

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ICS C23C004-12; C25B009-00
CC
     72-3 (Electrochemistry)
     Section cross-reference(s): 49
     7440-32-6, Titanium, uses
IT
        (electricity feeder made of Ti fiber-sintered plate whose surface
        is made smooth by plasma thermal spray of Ti in an
        inert gas (Ar, He, or N) atm.)
IT
     7440-37-1, Argon, uses
                              7440-59-7, Helium, uses 7727-37-9
     , Nitrogen, uses
        (electricity feeder made of a Ti fiber-sintered plate whose
        surface is made smooth by plasma thermal spray of Ti in
        an inert gas (Ar, He, or N) atm.)
IT
     7732-18-5, Water, processes
        (electricity feeders for water electrolytic cells)
IT
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
     Oxygen, preparation
        (electricity feeders for water electrolytic cells for prodn. of)
    ANSWER 9 OF 16 HCA COPYRIGHT 2006 ACS on STN
L63
125:333967 Direct solar-thermal hydrogen production from water using
     nozzle/skimmer and glow discharge. Pyle, W. R.;
    Hayes, M. H.; Spivak, A. L. (H-Ion Solar Co., Richmond, CA, 94805,
     USA). Proceedings of the Intersociety Energy Conversion Engineering
     Conference, 31st, 1753-1760 (English) 1996. CODEN:
     PIECDE. ISSN: 0146-955X. Publisher: Society of Automotive
     Engineers.
    An investigation of direct solar-thermal hydrogen and oxygen prodn.
AB
     from water is described. Nozzle jets and skimmers have been used
     for sepn. of the products and suppression of recombination.
     dissocn. of water vapor and the sepn. of its
     products was conducted in plasma-enhanced, non-equil.
     glow discharges.
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
IT
     Oxygen, preparation
        (direct solar-thermal hydrogen prodn. from water using
        nozzle/skimmer and glow discharge)
RN
     1333-74-0 HCA
    Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
    Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
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0 = 0

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

- CC 52-1 (Electrochemical, Radiational, and Thermal Energy Technology)
- ST solar thermal hydrogen prodn water; nozzle skimmer glow discharge hydrogen prodn
- IT Solar energy

(direct solar-thermal hydrogen prodn. from water using nozzle/skimmer and glow discharge)

IT Electric discharge

(glow, direct solar-thermal hydrogen prodn. from water using nozzle/skimmer and glow discharge)

IT Nozzles

(jet, direct solar-thermal hydrogen prodn. from water using nozzle/skimmer and **glow discharge**)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P,

Oxygen, preparation

(direct solar-thermal hydrogen prodn. from water using nozzle/skimmer and glow discharge)

IT 7732-18-5, Water, reactions

(direct solar-thermal hydrogen prodn. from water using nozzle/skimmer and glow discharge)

- L63 ANSWER 10 OF 16 HCA COPYRIGHT 2006 ACS on STN
- 123:68427 Development of vacuum plasma sprayed electrodes for an intermittently operated alkaline water electrolyser. Schiller, G.; Borck, V.; Henne, R.; Hug, W. (Deutsche Forschungsanstalt fur Luft- und Raumfahrt (DLR), Institute Technical Thermodynamics, Stuttgart, 70569, Germany). Hydrogen Energy Prog. X, Proc. World Hydrogen Energy Conf., 10th, Volume 1, 631-9. Editor(s): Block, David L.; Veziroglu, T. Nejat. Fla. Sol. Energy Cent.: Cape Canaveral, Fla. (English) 1994. CODEN: 61FXA7.
- AB Activated electrodes for advanced, intermittently operated alk. water electrolysis were developed by applying the vacuum plasma spraying technol. (VPS). For hydrogen evolution cathode coatings from Mo-contg. Raney nickel were used, and as anodes matrix composite coatings consisting of Raney nickel and Co3O4 electrocatalyst are applied. To test the electrodes for intermittent operation small electrolyzers with zero gap design were equipped with VPS electrodes of an area of 250 cm2 and NiO diaphragms. The electrochem. activity and the long-term degrdn. behavior of the electrodes were studied by applying const. c.d. as

well as a real solar c.d. profile. Polarization curve measurements in dependence of temp. were performed and single electrode potentials were detd. At std. conditions (300 mA/cm2, 80°) cell voltage was in the range of 1.6 V. During various operation modes with const. and intermittent power supply cell voltage was monitored up to 5000 h of operation showing stable behavior during the whole period of study without evidence of degrdn. The next step in the development of VPS electrodes is the prodn. of electrodes with an area of 600 cm2 which are currently prepd. to be tested in an intermittently operated 10 kW electrolyzer of advanced technol. 7732-18-5, Water, processes (development of vacuum plasma sprayed electrodes for intermittently operated alk. water electrolyzer) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 1333-74-0P, Hydrogen, properties 7782-44-7P, Oxygen, properties (development of vacuum plasma sprayed electrodes for intermittently operated alk. water electrolyzer) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0CC 72-2 (Electrochemistry) Section cross-reference(s): 49 vacuum plasma sprayed electrode water electrolysis Electrodes (vacuum **plasma** sprayed) 7732-18-5, Water, processes (development of vacuum plasma sprayed electrodes for intermittently operated alk. water electrolyzer) 1333-74-0P, Hydrogen, properties 7782-44-7P, Oxygen, properties (development of vacuum plasma sprayed electrodes for

intermittently operated alk. water electrolyzer)

ΙT

RN

CN

H<sub>2</sub>O

IT

RN

CN

H-H

RN

CN.

ST

IT

IT

IT

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ANSWER 11 OF 16 HCA COPYRIGHT 2006 ACS on STN
121:233994 Magnetohydrodynamic (MHD) process for the manufacture of
     hydrogen. Giannotti, Pedro (Brazil). Braz. Pedido PI BR 9204950 A
     19940614, 10 pp. (Portuguese). CODEN: BPXXDX.
     APPLICATION: BR 1992-4950 19921209.
AB
     The process comprises introducing water into a MHD reactor through a
     small bottom tube equipped with a needle valve and rotameter for
     flow control, ionizing the water with an electron beam in an elec.
     field to form a low-resistance elec. conductive plasma and
     sepg. the ions and electrons by elec. and magnetic forces to recover
     O at one side and H at the other side.
ΙT
     7732-18-5, Water, reactions
        (in MHD process for the manuf. of hydrogen)
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H<sub>2</sub>O
IT
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
     Oxygen, preparation
        (in MHD process for water dissocn)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
     7782-44-7 HCA
RN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
0 = 0
IC
     ICM C01B003-06
CC
     49-1 (Industrial Inorganic Chemicals)
     Section cross-reference(s): 77
     7732-18-5, Water, reactions
IT
        (in MHD process for the manuf. of hydrogen)
ΙT
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
     Oxygen, preparation
        (in MHD process for water dissocn)
     ANSWER 12 OF 16 HCA COPYRIGHT 2006 ACS on STN
L63
121:45138 Contact glow discharge electrolysis: a
     study of its chemical yields in aqueous inert-type electrolytes.
     Sengupta, Susanta K.; Singh, Om Prakash (Department of Chemistry,
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Faculty of Science, Banaras Hindu University, Varanasi, 221005.

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Journal of Electroanalytical Chemistry, 369(1-2), 113-20
     (English) 1994. CODEN: JECHES. ISSN: 0368-1874.
AΒ
     A study of the chem. yields of contact glow
     discharge electrolysis (CGDE) at the anode in aq. inert
     electrolytes at various quantities of electricity, applied voltages
     and electrolyte compns. shows that, for the passage of each mole of
     electrons, 0.25 mol of O2 and more than 1.0 mol of both H2 and H2O2
     plus O2 are produced at the anode when the glow
     discharges there are fully grown. Non-faradaic yields may
     originate in two reaction zones: the anolyte near the plasma
     where lig. water mols. are broken up into H2O2, O2 and H2, and the
     plasma around the anode where gas phase dissocn. of water
     mols. into H2 and O2 occurs. The former is important for anodic
           Hickling's radiolytic mechanism has been applied to interpret
     the chem. results of the liq. phase reaction zone. A comparative
     study of the chem. yields of anodic and cathodic CGDE indicates that
     the breakup of water mols. occurs entirely in the plasma
     during cathodic CGDE, but primarily in the liq. anolyte and partly
     in the plasma during the anodic phenomenon.
     7732-18-5, Water, reactions
IT
        (contact glow discharge electrolysis of, in
        aq. electrolytes)
RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H20
IT
     1333-74-0P, Hydrogen, preparation 7782-44-7P,
     Oxygen, preparation
        (evolution of, in contact glow discharge
        electrolysis of water)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
o = o
CC
     72-11 (Electrochemistry)
     Section cross-reference(s): 76
ST
     electrolysis glow discharge water; oxygen
     hydrogen peroxide formation water electrolysis; radiolysis Hickling
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mechanism water electrolysis ITRadiolysis (Hickling mechanism of, for water contact glow discharge electrolysis) Electrolysis IT (contact glow discharge, of aq. solns.) IT (in contact glow discharge electrolysis of water) IT Electric **discharge** (glow, in water electrolysis) 7778-80-5, Dipotassium sulfate, reactions IT (contact glow discharge electrolysis of aq., oxygen and hydrogen and hydrogen peroxide formation in) IT **7732-18-5**, Water, reactions (contact glow discharge electrolysis of, in aq. electrolytes) 7440-06-4, Platinum, uses IT (electrode, for contact glow discharge electrolysis of water) IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation (evolution of, in contact glow discharge electrolysis of water) IT 7722-84-1P, Hydrogen peroxide, preparation (formation of, in contact glow discharge electrolysis of water) ANSWER 13 OF 16 HCA COPYRIGHT 2006 ACS on STN 101:60956 Study of energetics and the mechanism of acoustic-chemical reactions. Relation of yields of hydrogen and hydrogen peroxide in different aqueous systems. Margulis, M. A.; Didenko, Yu. T. (Vses. Nauchno-Issled. Inst. Org. Sint., Moscow, USSR). Zhurnal Fizicheskoi Khimii, 58(6), 1402-5 (Russian) 1984. ISSN: 0044-4537. The formation kinetics of H2 in sonolysis of H2O in O, N, and air AB was studied and the initial rates of H2O2 formation under the effect of ultrasound on H2O in H atm. was detd. By using the exptl. data, the O concn. in cavitational bubble in the final stage of compression was calcd. It represents a condensed plasma of strongly compressed gas, with a small water vapor impurity, in supercrit. state. A mechanism is proposed, which explains the change of basic H2O-sonolysis products yield in N + O atm. as function of O concn. **7732-18-5**, reactions IT

(acoustic decompn. of, formation of hydrogen and hydrogen

peroxide in, oxygen concn. in bubbles during)

RN

7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT **7782-44-7P**, preparation

(formation and concn. of, in bubbles, during sonolysis of water)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IT 1333-74-0P, preparation

(formation of, in ultrasound decompn. of water, kinetics of)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

CC 67-3 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms)

Section cross-reference(s): 65

IT **7732-18-5**, reactions

(acoustic decompn. of, formation of hydrogen and hydrogen peroxide in, oxygen concn. in bubbles during)

IT **7782-44-7P**, preparation

(formation and concn. of, in bubbles, during sonolysis of water)

IT 1333-74-0P, preparation 7722-84-1P, preparation (formation of, in ultrasound decompn. of water, kinetics of)

L63 ANSWER 14 OF 16 HCA COPYRIGHT 2006 ACS on STN

96:71225 Thermal, electromagnetic system for producing hydrogen and oxygen. De Menezes, Maurilio (Brazil). Braz. Pedido PI BR 8001086 A 19810825, 5 pp. (Portuguese). CODEN: BPXXDX. APPLICATION: BR 1980-1086 19800225.

AB H and O are produced by thermal dissocn. of water with laser radiation, electron plasma, microwave radiation, or other means, in an app. having porous ceramic walls, which facilitate dissocn. and help prevent recombination in the low pressure atm.

IT **7732-18-5**, reactions

(decompn. of, by laser radiation heating, app. for)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H20 IT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by water thermal decompn., app. for) 1333-74-0 HCA RNHydrogen (8CI, 9CI) (CA INDEX NAME) CNH-HRN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IC C01B003-02; C01B013-02 49-1 (Industrial Inorganic Chemicals) . CC IT**7732-18-5**, reactions (decompn. of, by laser radiation heating, app. for) IT **1333-74-0P**, preparation **7782-44-7P**, preparation (manuf. of, by water thermal decompn., app. for) ANSWER 15 OF 16 HCA COPYRIGHT 2006 ACS on STN 95:177642 Solid polymer electrolyte water electrolysis. Takenaka, H.; Torikai, E.; Kawami, Y.; Wakabayashi, N. (Gov. Ind. Res. Inst., Ikeda, Japan). Advances in Hydrogen Energy, 2 (Hydrogen Energy Prog., Vol. 1), 107-17 (English) 1981. CODEN: AHENDB. ISSN: 0276-2412. AΒ Electrocatalysts and their plating on solid polymer electrolytes (SPE) were studied. Perfluorosulfonic acid polymer membranes (Nafion) were used as SPE. Noble metals and their alloys were directly attached to both sides of the membrane without a binder by special metal plating methods. The methods utilizing reactions of metal salt soln. with reducing agent on the membrane surface made it possible to increase the adherence of electrocatalysts to the membrane and also to eliminate almost totally the resistance of the electrocatalyst/SPE interface. Pretreatments of the membrane were also studied to improve the adherence. Hydrothermal and gas plasma treatments were more effective. The constituents of cell voltage, i.e., anodic and cathodic overvoltages and ohmic drop, were measured for 5 noble

metals and their alloys. The anodic overvoltage was a major

electrocatalysts. The anodic overvoltages increased in the

constituent of voltage losses and clearly depended on the kinds of

following order: Ir < Rh < Rh/Pt < Pt < Pt/Ru < Pd. Pure Ru had

high initial activity; however, it was corroded during O evolution. The electrodes based on Ir alloys were the best electrocatalysts for O evolution and had a Tafel slope of 40-60 mV/decade. The effects of operating temp. on cell performance were also studied. cathode and the anode were a thin layer of Pt and similar layer of Ir alloy, resp. The cell voltage decreased with an increase in temp. At c.d. 5 A/dm2 and at 90°, the cell voltage was 1.56-1.59 V corresponding to a thermal efficiency (based on  $\Delta$ H) of 93-5%. **7732-18-5**, reactions (electrolysis of, in cell with Nafion membrane and

IT

noble metal electrocatalysts)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT **1333-74-0P**, preparation

> (evolution of, in water electrolysis in cell with Nafion membrane and noble metal electrodes)

RN 1333-74-0 HCA

Hydrogen (8CI, 9CI) · (CA INDEX NAME) CN

H-H

IT **7782-44-7P**, preparation

> (evolution of, in water electrolysis in electrolytic cell with Nafion membrane and noble metal electrocatalysts)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

72-10 (Electrochemistry) CC Section cross-reference(s): 49, 67

solid polymer electrolyte water electrolysis; noble metal ST electrocatalyst water electrolysis; Nafion membrane electrocatalyst water electrolysis; etching Nafion membrane water electrolysis; oxygen evolution water electrolysis cell

Platinum-group metal alloys, base IT

Platinum-group metals

(anodes, on Nafion membrane for oxygen evolution in water electrolysis)

Plasma, chemical and physical effects IT (etching by, of Nafion membrane for oxygen evolution in water electrolysis)

IT Iridium alloy, base

(anodes, on Nafion membrane for oxygen evolution in water electrolysis)

TT 7439-88-5, uses and miscellaneous 7440-05-3, uses and miscellaneous 7440-06-4, uses and miscellaneous 7440-16-6, uses and miscellaneous

(anodes, on Nafion membrane for oxygen evolution in water electrolysis)

IT 11107-71-4 12779-05-4 39349-40-1

(anodes, on Nafion membrane for oxygen evolution in water electrolysis)

IT **7732-18-5**, reactions

(electrolysis of, in cell with Nafion membrane and noble metal electrocatalysts)

IT 1333-74-0P, preparation

(evolution of, in water electrolysis in cell with Nafion membrane and noble metal electrodes)

IT **7782-44-7P**, preparation

(evolution of, in water electrolysis in electrolytic cell with Nafion membrane and noble metal electrocatalysts)

IT 65506-90-3

(membrane, gas-plasma etching and hydrothermal treatment of, for electrocatalyst adherence in water electrolysis)

L63 ANSWER 16 OF 16 HCA COPYRIGHT 2006 ACS on STN

94:177482 Dissociation of water by microwave plasma.

(Akiyama, Morio, Japan; Tai, Kosuke; Inada, Takeshi). Jpn. Kokai Tokkyo Koho JP 56017902 19810220 Showa, 2 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1979-93457 19790720.

AB **H2O** is **vaporized** dropwise in an air or inert gas stream and dissocd. at .apprx.2500° in a **microwave** discharge **plasma** at ≥915 MHz around an anode surrounded by a perforated and cooled catalyst metal, the paramagnetic O and nonmagnetic H being projected in different directions.

IT **7732-18-5**, reactions

(dissocn. of, in microwave discharge)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H20

IT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by water dissocn. by microwave discharge)

RN 1333-74-0 HCA

Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN 0 = 0IC C01B003-04; C01B013-02; C25B005-00 CC 49-1 (Industrial Inorganic Chemicals) ST water dissocn microwave plasma; oxygen manuf water dissocn; hydrogen manuf water dissocn ΙT Microwave, chemical and physical effects (dissocn. by, of water for hydrogen and oxygen manuf.) IT **7732-18-5**, reactions (dissocn. of, in microwave discharge) **1333-74-0P**, preparation **7782-44-7P**, preparation IT (manuf. of, by water dissocn. by microwave discharge) => D L64 1-22 CBIB ABS HITSTR HITIND ANSWER 1 OF 22 HCA COPYRIGHT 2006 ACS on STN 144:174310 Integrated plasma fuel cell process. Steinberg, Meyer (HCE, L.L.C., USA). U.S. Pat. Appl. Publ. US 2006024538 A1 20060202, 8 pp., Cont.-in-part of U.S. Ser. No. 604,153, abandoned. (English). CODEN: USXXCO. APPLICATION: US 2003-604961 20030828. PRIORITY: US 2003-2003/604153 20030627. A method for efficiently producing energy, carbon, carbon monoxide, AΒ synthetic carbonaceous liq. and gaseous fuels and hydrogen from fossil or biomass fuels with minimal carbon dioxide emissions is disclosed. 7782-44-7P, Oxygen, preparation IT(integrated plasma fuel cell process) 7782-44-7 HCA RN CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IT 1333-74-0P, Hydrogen, uses (integrated plasma fuel cell process) RN 1333-74-0 HCA

Hydrogen (8CI, 9CI) (CA INDEX NAME)

CN

H-H

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INCL 429017000; 429019000; 429021000
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     Section cross-reference(s): 72
ST
     plasma fuel cell process integrated
IT
     Hydrocarbons, uses
        (C1-4; integrated plasma fuel cell process)
IT
     Thermodynamic cycle
        (Rankine, steam boiler; integrated plasma
        fuel cell process)
IT
     Water gas shift reaction
        (app.; integrated plasma fuel cell process)
     Fischer-Tropsch reaction
IT
        (catalytic reactor; integrated plasma fuel cell
        process)
     Fuel cells
IT
        (direct carbon; integrated plasma fuel cell process)
IT
     Ashes (residues)
     Diesel fuel
     Electrolytic cells
       Plasma
        (integrated plasma fuel cell process)
IT
     Gasoline
        (integrated plasma fuel cell process)
IT
     Carbonaceous materials (technological products)
        (integrated plasma fuel cell process)
IT
     Reactors
        (plasma; integrated plasma fuel cell process)
IT
     Fuel cells
        (solid oxide; integrated plasma fuel cell process)
IT
     7782-44-7P, Oxygen, preparation
        (integrated plasma fuel cell process)
IT
     67-56-1P, Methanol, uses 74-82-8P, Methane, uses
                                                           630-08-0P,
     Carbon monoxide, uses 1333-74-0P, Hydrogen, uses
     7440-44-0P, Carbon, uses 7704-34-9P, Sulfur, uses
        (integrated plasma fuel cell process)
    ANSWER 2 OF 22 HCA COPYRIGHT 2006 ACS on STN
L64
137:254703 Apparatus for producing electrical and heat energy, hydrogen,
     and oxygen. Kanarev, F. M.; Peisakhovich, Yu. A.; Podobedov, V. V.
```

AB App. for producing elec. and heat energy, hydrogen, and oxygen is described. The app. case made of insulating material has bottom lug and bottom lid both constituting electrode-to-electrode chamber

2177512 C1 20011227, No pp. given (Russian).

RUXXE7. APPLICATION: RU 2000-119666 20000724.

(Kubanskii Gosudarstvennyi Agrarnyi Universitet, Russia). Russ. RU

CODEN:

divided by bottom cylindrical lug into anode and cathode spaces. Flat annular anode has holes and is placed in anode space of electrode-to-electrode chamber. Rod-type cathode made of high-melting material is placed in insulating rod which is driven by means of its outer thread into threaded hole of bottom lug and centered in outlet pipe hollow. Outlet pipe and case form near-cathode space. Cylinder-shaped wound device for building magnetic field is fitted onto bottom lid so that cathode and near-cathode space are under impact of its magnetic field. Electrodes designed to take off elec. current generated are mounted in cylindrical part of anode space. Steam-gas mixt. outlet pipe is arranged in case concentrically relative to cathode, oxygen outlet pipes are mounted in top part of anode space. app. has enhanced power characteristics. 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes (app. for producing elec. and heat energy, hydrogen, and oxygen) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0**7732-18-5**, Water, reactions (electrolysis, high temp.; in app. for producing elec. and heat energy, hydrogen, and oxygen) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) ICM C25B001-02 ICS C25B009-00 72-9 (Electrochemistry) Section cross-reference(s): 47, 76 Decomposition (plasma; of water in app. for producing elec. and heat energy, hydrogen, and oxygen) 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes

(app. for producing elec. and heat energy, hydrogen, and oxygen)

IT

RN CN

H-H

RN

CN

IT

RN

CN

H<sub>2</sub>O

IC

CC

IT

IT

L64 ANSWER 3 OF 22 HCA COPYRIGHT 2006 ACS on STN

136:361000 Gear to generate thermal energy, hydrogen and oxygen.

Kanarev, F. M.; Konarev, V. V.; Podobedov, V. V. (Kubanskii
Gosudarstvennyi Agrarnyi Universitet, Russia). Russ. RU 2167958 C2

20010527, No pp. given (Russian). CODEN: RUXXE7.

APPLICATION: RU 1999-111975 19990602.

App. to generate thermal energy, hydrogen and oxygen is disclosed. AΒ App. has frame with axial hole and cylindrical-conical boss and lower cover which form jointly with frame interelectrode chamber including intercommunicating anode and cathode spaces. Circular anode with hole is positioned in anode space and rod cathode in put into dielec. rod which is brought into interelectrode chamber through threaded hole in lower cover. It enables working part of cathode to be centered relative to hole in refractory metal bushing put through axial hole of frame. Diam. of hole in bushing is less than diam. of cathode. Branch pipe to feed working soln. is located in anode chamber, branch pipes to release oxygen are brought to upper part of anode space, branch pipes to release hydrogen are brought to upper part of cathode space and branch pipe to remove steam and gas mixt. is installed in axial hole of frame. Given gear makes it feasible to generate thermal energy by heating of soln. with the use of plasma and to produce simultaneously hydrogen and oxygen by way of electrolytic and thermal decompn. of water.

IT 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes

(app. for generating thermal energy, hydrogen and oxygen)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-- H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

 $\circ = \circ$ 

IT 7732-18-5, Water, reactions

(electrolysis; app. for generating thermal energy, hydrogen and oxygen using).

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IC ICM C25B001-02 ICS C25B009-00

CC 72-9 (Electrochemistry)
 Section cross-reference(s): 47, 76

ST app generation heat hydrogen oxygen water electrolysis plasma dissocn

IT Dissociation

(plasma; app. for generating thermal energy, hydrogen and oxygen using)

IT 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes

(app. for generating thermal energy, hydrogen and oxygen)

L64 ANSWER 4 OF 22 HCA COPYRIGHT 2006 ACS on STN 136:360987 Electrochemical micro gap/micro-channel electrolysis. Schlueter, Claus (Germany). Ger. Offen. DE 10054157 A1 20020508, 6 pp. (German). CODEN: GWXXBX. APPLICATION: DE 2000-10054157 20001102.

Decompn. of water as hydrogen energy storage and renewable energy AB was used in underpotential electrochem. micro gap/micro-channel electrolytic cells operated using the electrochem. voltage gradient on selected metals. Electrochem. micro gap/micro-channel electrolysis under the underpotential using selected metals in the micro gap/micro-channel electrolytic liq. under high current led without creation of an external potential was used as renewable hydrogen prodn. system. Two-stage electrolytic/chem. procedure of the electrolytic decompn. from NaOH to Na and OH and chem. reaction of Na with water for NaOH with hydrogen evolution was used. light-wt., wide-spread sheet metal foil and fine grid sandwich construction was employed in continuous economic manufq. with the increased efficiency using the electrolyzer. Waste heat cooling/exhaust gas were used in gas turbine cooling systems. or H/O high pressure gas injection plasma combustion chamber with high temp.-firm W or Ta glowing elements was employed.

IT 7732-18-5, Water, reactions

(electrochem. micro gap/micro-channel electrolysis)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation (electrochem. micro gap/micro-channel electrolysis) RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IC ICM C25B001-00 72-3 (Electrochemistry) CC Section cross-reference(s): 49, 52 IT 1310-73-2, Sodium hydroxide, reactions **7732-18-5**, Water, reactions (electrochem. micro gap/micro-channel electrolysis) IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation (electrochem. micro gap/micro-channel electrolysis) ANSWER 5 OF 22 HCA COPYRIGHT 2006 ACS on STN 136:56365 Energy conversion process and device. Artamonov, A. S. (Sergeevich, Artamonov Aleksandr, Russia). Russ. RU 2154738 C2 20000820, No pp. qiven (Russian). CODEN: RUXXE7. APPLICATION: RU 1997-121735 19971209. An elec.-conducting lig. undergoes electrothermal dissocn. with a AB lig. or solid hydrocarbon fuel introduced into a heating zone wherein the elec.-conducting lig. is injected. Heating and evapn. processes involve burst of injected sprays by periodic excitation of elec. discharges in them to produce a vapor-gas mixt. or H2+O2, or a mixt. of disintegration products of the solid fuel and liq. An app. implementing the process contains reactors with covers, mixing chambers, a steam collector, a steam turbine, an elec. generator, a rotary elec.-pulse generator, a transformer, a surface condenser with a condensate purifn. unit, and a chamber for

prepg. an electrolyte and superheated water. The app. is also provided with burners arranged in the reactor covers, condensate injectors, addnl. reactors, and mixing chambers. Each burner has injectors with pipe connections incorporating cylindrical endless screw feeders and central electrodes in the form of cylindrical

chambers' connected to the nozzle and covered with an elec. insulator on outer surfaces. Another design version of the app. has an engine

contg. ≥1 cylinder with a piston, a crank gear coupled with a crankshaft, a combustion chamber, a transportation system for the soln. and fuel, an air supply system, a waste gas exhaust systems, and an elec. charge excitation system. The engine is provided with a composite injector with a pipe connection to the endless screw feeder, a central electrode, a fuel injector, and a water injector for the combustion chamber. Each engine cylinder is provided with addnl. combustion chambers and a fuel feed system. The crank gear consists of 2 parts braced together by means of a spring and an anchor bolt for their relative sliding.

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation

(formation in energy conversion process)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IC ICM F01K027-00

ICS F02B045-10; F01D015-00

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 47

IT Plasma

(in energy conversion process)

IT 1333-74-0P, Hydrogen, preparation 7782-44-7P,
Oxygen, preparation
 (formation in energy conversion process)

L64 ANSWER 6 OF 22 HCA COPYRIGHT 2006 ACS on STN

135:155255 Water-fueled engines, method and system for power generation, and machine systems. Hatanaka, Takeshi (Japan). Jpn. Kokai Tokkyo Koho JP 2001221060 A2 20010817, 9 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-67694 20000207.

AB The engine comprises (a) a means for feeding water equipped with a no-return valve, (b) a means for generation of hydrogen-oxygen gas from water by plasma arc treatment,

(d) a means for generation of high-temp. and -pressure **steam** by reaction of the gas, and (e) a means for generation of power from the **steam**. Also claimed are (A) method and a system for storage of the generated power as rotational energy, in flywheel turbines and (B) a machine system comprising the engines and

optionally a motor. The engines and the systems are environmentally friendly. IT 1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen, uses (engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) 1333-74-0 RN HCACN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN o = 0IC ICM F02C003-28 B60K003-04; F01D001-04; F01D005-12; F02C006-00; B01J019-08; ICS B60K006-02 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC environmentally friendly water fueled engine; motor plasma ST treated water fuel engine; hydrogen oxygen fuel environmentally friendly engine IT Electric motors Engines Plasma Seawater (engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) IT Turbines (flywheel; engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) IT (generation; engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) 1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen, IT uses (engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) 7722-84-1P, Hydrogen peroxide, uses IT (water contg.; engines operated by hydrogen-oxygen fuel generated by plasma treatment of water) ANSWER 7 OF 22 HCA COPYRIGHT 2006 ACS on STN L64

135:155249 Closed-cycle hydrogen engines, system and method for power

generation, and machine systems. Hatanaka, Takeshi (Japan). Kokai Tokkyo Koho JP 2001221013 A2 20010817, 10 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-67695 20000207. The closed-cycle hydrogen engine comprises (a) an expansion turbine, AB (b) a high-pressure pump for pressurization of electrolyte solns., (c) a means for generation of mixt. gas of hydrogen-oxygen and steam from electrolyte soln. by plasma arc treatment, (d) a means for combustion of the mixt. gas for generation of high-temp. and -pressure steam, (e) a jet nozzle for feeding the high-temp. and -pressure steam to the turbine, (f) a condenser for generation of electrolyte soln. from the turbine waste gas, and (g) a circulation cycle connecting the condenser and the pump. Also claimed are (A) the above stated engines placed in an engine housing, (B) method for power generation with the engine, (C) a system for the power generation, and (D) a machine system comprising the engines and optionally a motor. engines and the systems are environmentally friendly. 1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen, IT uses (fuel; closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-H7782-44-7 HCA RN Oxygen (8CI, 9CI) (CA INDEX NAME) CN o = oIC ICM F01K025-00 ICS B60K006-02; C01B003-02; C01B003-04; F01D005-02; F01D009-02; F01K015-02; F02C003-22 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC ST environmentally friendly closed system hydrogen engine; oxygen hydrogen fuel environmentally friendly engine; motor closed system hydrogen engine; electrolyte circulation closed system turbine engine; plasma arc electrolyte conversion fuel engine IT Inks (India; closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors) IT Condensers Electric motors Electrolytes

Engines

Plasma

Seawater

Turbines

Waters

(closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors)

IT Power

(generation; closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors)

IT 7722-84-1P, Hydrogen peroxide, uses

(electrolyte; closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors)

IT 1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen, uses

(fuel; closed-cycle hydrogen engines operated by fuel generated from electrolyte solns. in plasma reactors)

L64 ANSWER 8 OF 22 HCA COPYRIGHT 2006 ACS on STN

134:165236 Method and apparatus for manufacture of hydrogen, and power system using same.. Hatanaka, Takeshi (Japan). Jpn. Kokai Tokkyo Koho JP 2001039701 A2 20010213, 8 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1999-243651 19990728.

The method comprises arranging multiphase a.c. electrodes in a AB casing for forming plural reaction zones along those electrodes, supplying multiphase a.c. power to the multiphase a.c. electrodes for generating plasma arcs in the plural reaction zones in order, supplying water to the reaction zones to generate steam, and catalytically reacting that in the plasma arcs of the reaction zones in order to obtain H2 and O2. includes micro-reaction-path means packed in the reaction zones, and a neutral electrode arranged opposite to the multiphase a.c. electrodes in the casing; the neutral electrode is connected to the neutral point of the multiphase a.c. power. The power system includes a water supply source, a H2 manuf. app. for generating a H2- and O2-contg. fuel by decompn. of water, a prime mover driven by combustion of the fuel to generate mech. output, and a power source driven by the prime mover.

IT 7732-18-5, Water, reactions

(decompn. of; method and app. for manuf. of hydrogen and power system using same)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0P, Hydrogen, preparation (fuel contq.; method and app. for manuf. of hydrogen and power

system using same)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IC ICM C01B003-04 ICS F22B001-30

CC 49-1 (Industrial Inorganic Chemicals)
Section cross-reference(s): 52

ST hydrogen manuf app **plasma** water decompn; power system hydrogen manuf app

IT Plasma

(method and app. for manuf. of hydrogen and power system using same)

IT 1333-74-0P, Hydrogen, preparation (fuel contg.; method and app. for manuf. of hydrogen and power system using same)

L64 ANSWER 9 OF 22 HCA COPYRIGHT 2006 ACS on STN

128:129590 Plasma burner for reactive decomposition of gas streams. Preuschen, Erwin G. (Preuschen, Erwin G., Germany). Ger. Offen. DE 19628932 Al 19980129, 6 pp. (German). CODEN: GWXXBX. APPLICATION: DE 1996-19628932 19960718.

AB A plasma burner is described for sep. of a water vapor stream into H2 and O2. The water vapor stream is injected through a lance in the anode bore of the burner comprising a coaxial cathode/anode unit. The burner can be heated by a nuclear reactor. The housing downstream of the burner contains devices for generating an electrostatic field at the discharge openings which also contain mol. grids from Pt or Pt alloys or Pt-coated porous ceramics. The sepd. H2 can be

recirculated to the plasma burner. ΙT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation (plasma burner for sepg. water vapor streams into hydrogen and oxygen) RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN 0 = 07732-18-5, Water, reactions. IT (plasma burner for sepg. water vapor streams into hydrogen and oxygen) RN 7732-18-5 HCA CNWater (8CI, 9CI) (CA INDEX NAME) H<sub>2</sub>O IC ICM B01D057-00 ICS B01D053-00; C01B003-50 47-3 (Apparatus and Plant Equipment) CC ST plasma burner water sepn hydrogen oxygen IT Anodes Burners Cathodes Electrostatic field Lances Thermal decomposition (plasma burner for sepg. water vapor streams into hydrogen and oxygen) IT Platinum alloy (plasma burner for sepg. water vapor streams into hydrogen and oxygen) IT 7440-06-4, Platinum, uses (plasma burner for sepg. water vapor streams into hydrogen and oxygen) 1333-74-0P, Hydrogen, preparation 7782-44-7P, IT Oxygen, preparation (plasma burner for sepg. water vapor streams into hydrogen and oxygen)

IT **7732-18-5**, Water, reactions (plasma burner for sepg. water vapor streams into hydrogen and oxygen) ANSWER 10 OF 22 HCA COPYRIGHT 2006 ACS on STN L64 125:343576 Plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber. Yamazaki, Shunpei (Handotai Energy Kenkyusho, Japan). Jpn. Kokai Tokkyo Koho JP 08241869 A2 **19960917** Heisei, 8 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1996-61890 19960223. Impurity in a plasma reaction chamber, used for forming a AB film, is removed by converting H and HCl or Cl to plasma. The method removes an impurity such as O and alkali metals from inner wall of the chamber which is used for manuf. of a p-type or n-type semiconductor layer and a photoconductor. IT 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes (removal; in plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber) RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IC ICM H01L021-205 ICS C23C016-50; C23F004-00; H01L021-3065 ICA C23C014-00 CC 75-1 (Crystallography and Liquid Crystals) Section cross-reference(s): 76 semiconductor manuf plasma CVD chamber cleaning; ST photoconductor manuf plasma CVD chamber cleaning Plasma ΙT (formation; plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber) ITSemiconductor devices (manuf.; plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber) IT Electric conductors

(photoconductive; plasma treatment for removal of impurity attached on inner wall of plasma reaction

chamber)

- IT Water vapor
  - (removal; in plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber)
- IT Alkali metals, processes
  - (removal; in plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber)
- IT Vapor deposition processes
  - (plasma, plasma treatment for removal of impurity attached on inner wall of plasma reaction chamber)
- IT 993-07-7, Trimethylsilane 7803-62-5, Silane, reactions
  (plasma CVD film raw material; in plasma
  treatment for removal of impurity attached on inner wall of
  plasma reaction chamber)

- IT 12586-59-3P, Proton
  (plasma; plasma treatment for removal of
  - impurity attached on inner wall of plasma reaction chamber)
- IT 1333-74-0P, Hydrogen, processes 7782-44-7P, Oxygen, processes (removal; in plasma treatment for removal o
  - (removal; in **plasma** treatment for removal of impurity attached on inner wall of **plasma** reaction chamber)
- L64 ANSWER 11 OF 22 HCA COPYRIGHT 2006 ACS on STN
- 123:269320 Vacuum plasma spraying of high-performance electrodes for alkaline water electrolysis. Schiller, G.; Henne, R.; Borck, V. (Institute of Technical Thermodynamics, Deutsche Forschungsanstalt Luft- und Raumfahrt, Stuttgart, 70569, Germany). Journal of Thermal Spray Technology, 4(2), 185-94 (English) 1995. CODEN: JTTEE5. ISSN: 1059-9630.
- AB Electrode coatings for advanced alk. water electrolysis were produced by applying the vacuum plasma spraying (VPS) process. The characteristics of the used VPS equipment that were essential for the development of effective electrocatalytic electrode layers are presented. Mo-contg. Raney Ni coatings were applied for cathodic H2 evolution, and Raney Ni/Co3O4 matrix composite layers were developed for the anodic O2 evolution reaction. For the prepn. of Raney Ni coatings, a precursor alloy such as Ni-Al was sprayed that had to be leached subsequently in caustic soln. to remove the Al content, forming a porous,

IT

RN

CN -

H-H

IT

RN CN

IT

RN

CN

H<sub>2</sub>O

CC

ST

IT

IT

IT

Spraying

high-surface-area Ni layer. The spray powders and the resulting VPS layers were studied by metallog., x-ray diffraction (XRD), and SEM/energy dispersive anal. by x-ray(EDX). For spraying of thermally sensitive oxide electrocatalysts (e.g., Co3O4), special process conditions involving plasma-chem. effects (reactive plasma spraying) had to be developed. The electrocatalytic activity of the electrode coatings was investigated by performing polarization curves free of ohmic losses (IR-free) and long-term tests under conditions of continuous and intermittent operation, which showed excellent electrochem. properties. 1333-74-0P, Hydrogen, preparation (evolution of hydrogen in alk. water electrolysis using nickel electrodes) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7782-44-7P, Oxygen, preparation (evolution of oxygen in alk. water electrolysis using nickel electrodes) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) o = 0**7732-18-5**, Water, reactions (vacuum plasma spraying of nickel on high-performance electrodes for alk. water electrolysis) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 72-9 (Electrochemistry) Section cross-reference(s): 52, 56 vacuum plasma spraying electrode water electrolysis; alk water electrolysis high performance electrode Electrolytic polarization (polarization of vacuum plasma sprayed Raney nickel electrodes in alk. water electrolysis) Electrodes (vacuum plasma spraying of high-performance electrodes for alk. water electrolysis)

(vacuum plasma spraying of nickel and nickel alloy electrodes for hydrogen and oxygen manuf. from alk. water electrolysis)

IT 164850-48-0

(Al-Mo-Ni alloy precursor for producing vacuum plasma sprayed Raney nickel electrodes in alk. water electrolysis)

- IT 1333-74-0P, Hydrogen, preparation (evolution of hydrogen in alk. water electrolysis using nickel electrodes)

- IT 1310-58-3, Potassium hydroxide, uses
   (vacuum plasma spraying of nickel on high-performance
   electrodes for alk. water electrolysis)
- TT 7732-18-5, Water, reactions
   (vacuum plasma spraying of nickel on high-performance
   electrodes for alk. water electrolysis)
- L64 ANSWER 12 OF 22 HCA COPYRIGHT 2006 ACS on STN

  122:121076 Plasma ashing apparatus. Sasaki, Takashi; Harada,
  Michuki; Hitomi, Shuji; Yoneyama, Shimao (Shinko Pantec Co Ltd,
  Japan; Mitsubishi Corp; Japan Storage Battery Co Ltd; Emu Shii
  Erekutoronikusu Kk). Jpn. Kokai Tokkyo Koho JP 06151386 A2

  19940531 Heisei, 7 pp. (Japanese). CODEN: JKXXAF.
  APPLICATION: JP 1992-295058 19921104.
- AB In manuf. of plasma ashing app. comprising H2 and O2 gas sources and a plasma asher for removing resist used in semiconductor product processes by plasma generated in an ashing chamber using O2 gas, mixt. of O2 gas and H2 gas or water vapor formed by mixing O2 gas and H2 gas from the gas sources, the gas source is a water electrolysis device.

IT 1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen, uses

(manuf. of plasma ashing app.)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

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RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
0 = 0
IT
     7732-18-5, Water, uses
        (manuf. of plasma ashing app.)
RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
IC
     ICM
          H01L021-302
     ICS H01L021-027
CC
     76-14 (Electric Phenomena)
ST
     plasma ashing app water electrolysis device; oxygen
     hydrogen water electrolysis device
IT
     Resists
        (manuf. of plasma ashing app. for removing of)
IT
     Ashing
        (plasma, manuf. of plasma ashing app.)
IT
     1333-74-0P, Hydrogen, uses 7782-44-7P, Oxygen,
        (manuf. of plasma ashing app.)
     7732-18-5, Water, uses
IT
        (manuf. of plasma ashing app.)
     ANSWER 13 OF 22 HCA COPYRIGHT 2006 ACS on STN
117:57606 Vacuum plasma sprayed electrodes for advanced
     alkaline water electrolysis. Schiller, G.; Borck, V. (Inst. Tech.
     Thermodyn., Ger. Aerosp. Res. Estab., Stuttgart, 7000/80, Germany).
     International Journal of Hydrogen Energy, 17(4), 261-73 (English)
            CODEN: IJHEDX.
                            ISSN: 0360-3199.
     1992.
     Electrode coatings for advanced alk. water electrolysis were
AΒ
     produced by applying vacuum plasma spraying (VPS). The
     characteristics of the VPS-equipment used which are essential for
     developing effective electrocatalytic electrode layers are
     presented.
                Raney nickel and Raney nickel/Mo coatings were applied
     for cathodic hydrogen evolution whereas Co3O4 spinel and Raney
     nickel/Co304 composite layers served as electrocatalytic coatings
     for anodic oxygen evolution. The spray powders and the VPS layers
     were studied by x-ray diffraction and high resoln. SEM.
     surface prepn. of the substrates is required to achieve well-bonded
     and stable layers esp. with NiAl precursor alloy which has to be
     activated subsequently to Raney nickel. Due to the fact that oxide
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electrocatalysts such as Co304, for example, exhibit a high tendency

for oxygen loss and decompn. and, thus, loss of electrocatalytic activity at elevated temps. special process conditions and parameters had to be developed. "Reactive plasma spraying" involving plasmachem. effects allows the processing of thermal sensitive oxides to stable and undecomposed layers. The electrocatalytic activity of the electrode coatings was investigated by performing IR-free polarization curves up to current densities of 1 A cm-1 and long-time tests under continuous current loading of 0.5 A cm-2 over a period of 3000 h. The cathodic layers were found to exhibit overvoltages of 70-90 mV at A cm-2 and 70° in 25% KOH soln. Raney nickel/Co3O4 composite anodes, which are still in the initial state of development, show overvoltage values of 290 mV at 1 A cm-2.

IT **7732-18-5**, Water, reactions

(electrolysis of alk., vacuum plasma spray electrode for)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H20

IT 7782-44-7P, Oxygen, preparation

(evolution of, anodic, Raney nickel/cobalt oxide composite layers as electrocatalytic coating for)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IT 1333-74-0P, Hydrogen, preparation

(evolution of, on different Raney nickel coating, compared with uncoated nickel)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

CC 72-2 (Electrochemistry)

Section cross-reference(s): 49, 56

ST vacuum plasma sprayed electrode water electrolysis; Raney nickel molybdenum coated electrode; cobalt oxide Raney nickel composite electrode

IT Surface structure

(of Raney nickel plasma-sprayed coatings, for cathodes)

IT 7732-18-5, Water, reactions

(electrolysis of alk., vacuum plasma spray electrode for)

- IT 7782-44-7P, Oxygen, preparation
   (evolution of, anodic, Raney nickel/cobalt oxide composite layers
   as electrocatalytic coating for)
- 1333-74-0P, Hydrogen, preparation
   (evolution of, on different Raney nickel coating, compared with
   uncoated nickel)
- IT 12004-71-6 12043-39-9 12635-27-7 (vacuum plasma spraying of, activated subsequently to Raney nickel, for alk. water electrolysis)
- L64 ANSWER 14 OF 22 HCA COPYRIGHT 2006 ACS on STN
  116:160736 Water radiolysis under NET conditions. Lorenzetto, P.;
  Bjergbakke, E.; Hickel, B. (NET Team, IPP, Garching, D-8046,
  Germany). Fusion Engineering and Design, 17, 265-70 (English)
  1991. CODEN: FEDEEE. ISSN: 0920-3796.
- Radiolytic decompn. of the NET plasma facing component and AB . blanket water coolant is expected due to the intense radiation fields present inside the vacuum vessel. A sensitivity anal. was performed by computer simulation to study the formation and the suppression of the radiolytic products under NET conditions. formation of the stable radiolytic products (oxygen, hydrogen and hydrogen peroxide) formed within the water due to the decompn. induced by mixed neutron and gamma irradn. is given for several cases, which are relevant to the first wall water coolant. influence of coolant temp. (20, 70, 130, and 200°C), heterogeneous decompn. of H2O2, added solutes such as hydrogen, and ferrous ions simulating possible corrosion products was studied. There are still large discrepancies in the literature on the radiation chem. yields (G-values) in the case of the neutron irradn. Therefore 3 sets of G-values were considered, and the sensitivity of the applied G-values for mixed neutron and gamma irradn. was investigated. For all these cases, the hydrogen concn. required to suppress the water radiolysis is given, and some crit. issues with respect to the specific contribution of neutrons with high energy typical for the fusion spectrum (10-14 MeV) on the radiolytic water decompn. are discussed.
- IT 1333-74-0P, Hydrogen, preparation 7782-44-7P, Oxygen, preparation

(formation of, in radiolysis of water under tokamak conditions)

- RN 1333-74-0 HCA
- CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

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н—н
     7782-44-7 HCA
RN
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
o = o
IT
     7732-18-5, Water, reactions
        (radiolysis of, under tokamak conditions, hydrogen suppression
        of)
     7732-18-5 HCA
RN
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H<sub>2</sub>O
     71-2 (Nuclear Technology)
CC
     Section cross-reference(s): 74
IT
     Nuclear fusion reactors, tokamak
        (plasma-facing component of, radiolytic decompn. of
        coolant water in)
     1333-74-0P, Hydrogen, preparation 7722-84-1P, Hydrogen
IT
     peroxide, preparation 7782-44-7P, Oxygen, preparation
        (formation of, in radiolysis of water under tokamak conditions)
     7732-18-5, Water, reactions
IT
        (radiolysis of, under tokamak conditions, hydrogen suppression
        of)
    ANSWER 15 OF 22 HCA COPYRIGHT 2006 ACS on STN
106:164339 Electrocatalysts for advanced electrolysis of water.
     Schurnberger, W.; Divisek, J. (Stuttgart, Fed. Rep. Ger.).
     VDI-Berichte, 602, 63-78 (German) 1987. CODEN: VDIBAP.
     ISSN: 0083-5560.
AB
     A review with 19 refs.
                             The important catalysts for the prodn. of H
     and O are discussed. Among these are Raney Ni, spinels, perovskites
     and porous sintered Ni (or Ni alloys). The possibility of coating
     electrodes via plasma spraying is also considered.
     7732-18-5, Water, reactions
IT
        (electrolysis of, catalysts for)
     7732-18-5 HCA
RN
```

CN

Water (8CI, 9CI) (CA INDEX NAME)

IT 1333-74-0P, Hydrogen; preparation 7782-44-7P, Oxygen, preparation (prodn. of, in water electrolysis, catalysts for) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-H RN 7782-44-7 HCA. CN Oxygen (8CI, 9CI) (CA INDEX NAME)  $0 \stackrel{\cdot}{=} 0$ CC 72-0 (Electrochemistry)

- L64 ANSWER 16 OF 22 HCA COPYRIGHT 2006 ACS on STN

  101:237292 Low pressure plasma sprayed electrode coatings for alkaline water electrolysis. Schnurnberger, W.; Henne, R.; Von Bradke, M. (Inst. Tech. Phys., DFVLR, Stuttgart, 7000/80, Fed. Rep. Ger.). Advances in Hydrogen Energy, 4 (Hydrogen Energy Prog. 5, Vol. 2), 933-41 (English) 1984. CODEN: AHENDB. ISSN: 0276-2412.
- Electrodes coated with stable and active electrocatalysts allow H AB prodn. in alk. solns. at high c.d. and efficiency. Porous Raney Ni and mixed metal oxides (spinels, perovskites) used as electrode coatings reduce significantly the overvoltage of H2O decompn. By means of the low pressure plasma spraying method (LPPS), various electrocatalytic materials can be applied reliably without undue overheating of the substrates. Furthermore, in consequence of the short time needed for the LPPS coating process, thermal decompn. and structural changes of the catalysts can be largely avoided. The prepn. and properties of Raney Ni and Ni/Co-spinel coatings on Ni substrates are reported. The structure of the electrode coatings was studied by SEM and x-ray diffraction, indicating no significant change of the electrocatalysts subsequent to LPPS. The electrochem. behavior of the coatings was investigated with regard to evolution of both O and H in 25 wt. % KOH solns. at c.d. ≤1 A/cm2. Tafel-plots at 23°-100° are given. The LPPS coating technique can be used to activate both anodes and cathodes for electrolysis of alk. H2O.

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IT
     7732-18-5, reactions
        (electrolysis of, plasma-sprayed electrodes for)
     7732-18-5 HCA
RN
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H20
IT
     1333-74-0P, preparation 7782-44-7P, preparation
        (prodn. of, by water electrolysis, plasma-
        sprayed electrodes for)
RN '
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
CC
     72-9 (Electrochemistry)
     plasma sprayed electrode water
ST
     electrolysis; nickel Raney cobalt spinel electrode; hydrogen oxygen
     electroprodn water electrolysis
IT
     Electrodes
        (plasma-sprayed, for water
        electrolysis)
IT
     Coating process
        (plasma, on electrodes for water electrolysis)
ΙT
     12017-35-5
        (electrodes with coatings from plasma-sprayed
        , for water electrolysis)
     7440-02-0, uses and miscellaneous
IT
        (electrodes, plasma-sprayed, for
        water electrolysis)
     7732-18-5, reactions
IT
        (electrolysis of, plasma-sprayed electrodes for)
     1333-74-0P, preparation 7782-44-7P, preparation
IT
        (prodn. of, by water electrolysis, plasma-
        sprayed electrodes for)
L64 ANSWER 17 OF 22 HCA COPYRIGHT 2006 ACS on STN
100:182188 Developments in advanced alkaline water electrolysis. Bowen,
     C. T.; Davis, H. J.; Henshaw, B. F.; Lachance, R.; LeRoy, R. L.;
     Renaud, R. (Noranda Res. Cent., Pointe-Claire, QC, H9R 1G5, Can.).
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International Journal of Hydrogen Energy, 9(1-2), 59-66 (English) CODEN: IJHEDX. ISSN: 0360-3199. AB Two parallel alk. water electrolysis programs are underway at the Noranda Research Center. One of these is part of Canada's contribution to the International Energy Agency (IEA) H program. Its objective is the evaluation of new anode, cathode and separator materials on an industrial scale. Porous Teflon cloth impregnated with K titanate was shown to have the lowest resistance factor of the separators tested, but its fragility and hydrophobicity are of concern. Felted polysulfone has the strength, gas tightness and low resistivity required. For all separators, effective resistivity depended on the free space allowed, the strength of this effect increasing with separator hydrophobicity. Ni electrodes, plasma-sprayed with Ni/Al or Ni/stainless steel powders, gave electrocatalytic activities with high stability. program, development of Electrolyzer Inc.'s advanced unipolar electrolysis technol., has benefited from the IEA activity. obtained with 3 100-kA cells on the Generation I design after more than 1000 h of operation are presented, together with results obtained in theGeneration II configuration over a 5000 h period. IT **7732-18-5P**, preparation (electrolysis of alk., cells for) RN 7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME) H20 IT **1333-74-0P**, preparation **7782-44-7P**, preparation (prodn. of, in alk. water electrolysis, cells for) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-HRN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) o = 0CC 72-9 (Electrochemistry) Section cross-reference(s): 52 IT **7732-18-5P**, preparation (electrolysis of alk., cells for) IT 1333-74-0P, preparation 7782-44-7P, preparation

(prodn. of, in alk. water electrolysis, cells for)

L64 ANSWER 18 OF 22 HCA COPYRIGHT 2006 ACS on STN 97:225561 Anodic and cathodic catalysts for high-temperature high current density alkaline water electrolysis. Prigent, M.; Nenner, T. (Tech. Appl. Energet., Inst. Francais Petrole, Rueil Malmaison, 92506, Fr.). Advances in Hydrogen Energy, 3 (Hydrogen Energy Prog. 4, Vol. 1), 299-307 (English) 1982. CODEN: AHENDB. ISSN: 0276-2412.

AB Electrocatalysts for electrolysis of alk. H2O at a c.d. of 10 kA/m2 and a temp. of 160° are prepd. on a Raney Ni base prepd. by plasma gun spraying of a Ni-Al alloy on a Ni grid, followed by KOH leaching. X rays studies show that resistance to sintering at elevated temp. of the finely divided layer of Ni is improved by incorporating small quantities of a 3rd element like Ti, Zr, Mo. Evaluation of the catalyst is performed at 10 kA/m2, in 40% wt. KOH at 160°, the electrodes being pressed against the diaphragm. These electrodes used as cathode will give a voltage redn. of .apprx.210 mV compared to the non-activated electrode. These Raney electrodes covered with mixed oxide of Co, Sr, La give a stable redn. of the O overvoltage of 180 mV at 160°. When used in a couple, a total cell voltage of 1.75 V can be obtained at 160°.

IT **7732-18-5**, reactions

(electrolysis of alk., Raney nickel catalytic electrodes for)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by electrolysis of alk. water, Raney nickel catalysts for)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

CC 72-9 (Electrochemistry)
Section cross-reference(s): 49, 52, 67

IT **7732-18-5**, reactions

(electrolysis of alk., Raney nickel catalytic electrodes for)
IT 1333-74-0P, preparation 7782-44-7P, preparation
 (manuf. of, by electrolysis of alk. water, Raney nickel catalysts for)

L64 ANSWER 19 OF 22 HCA COPYRIGHT 2006 ACS on STN
91:65203 Electrolytic production of hydrogen from alkaline solutions active electrodes for potassium hydroxide electrolysis. Nidola, A.;
 Spaziante, P. M.; Giuffre, L. (Oronzio De Nora Impianti Elettrochim.
 S.p.A., Milan, Italy). Comm. Eur. Communities, [Rep.] EUR, EUR
 6085, Semin. Hydrogen Energy Vector: Prod., Use, Transp., 212-28
 (English) 1978. CODEN: CECED9.

New electrocatalysts for H and O evolution electrode reactions in AB KOH 29 wt. % electrolysis at atm. pressure and at 70° were studied and developed by Oronzio De Nora-Impianti Elettrochimici S.p.A.. Cathodic catalysts include Ru and Re deposited on Fe electrode bases via electrolytic or plasma jet techniques, while anodic catalysts include steel activated Ni and/or Co on Ni coated Fe or Ni electrode bases via the plasma jet route. The coatings with respect to com. catalysts, which include Ni and/or Fe sulfide, have the following practical advantages: low overvoltage for the desired reaction even at high c.d. (10 kA/m2), absence of catalytic aging phenomena vs. operation time and c.d., and no asbestos diaphragm deterioration. The relatively low amt. of catalysts (≤50 g/m2 for anodic coatings, ≤20 g/m2 for cathodic coatings) deposited on electrode bases makes them attractive for H prodn. by electrolysis of KOH solns. provided that a c. d. at least ≥3.0 kA/m2 is used.

IT **7732-18-5**, reactions

(electrolysis of, dissocn. catalysts in)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by electrolysis of water, catalysts in)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0CC 72-10 (Electrochemistry) Section cross-reference(s): 49, 67 7732-18-5, reactions IT (electrolysis of, dissocn. catalysts in) IT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by electrolysis of water, catalysts in) ANSWER 20 OF 22 HCA COPYRIGHT 2006 ACS on STN 91:11326 Electrochemical reduction of water vapor. Developmental stage of an assembly of planar cells. Viguie, J. C. (Cent. Etud. Nucl. Grenoble, Commis. Energ. At., Grenoble, Fr.). Comm. Eur. Communities, [Rep.] EUR, EUR 6085, Semin. Hydrogen Energy Vector: Prod., Use, Transp., 310-27 (French) 1978. CODEN: CECED9. AB The operating conditions of a planar electrolytic cell are described. The 50-cm2 half-cathodic cell was fabricated with a solid electrolyte of ZrO2-9 mol % Y2O3, assocd. with an embedded layer of Ni powder as the cathode. The disk is supported by a honeycomb-shaped ceramic covered by a layer of Ni. The solid electrolyte can be prepd. by powder compaction and sintering or by plasma jet projection. IT7732-18-5, vapor (electrochem. redn. of, in planar cell) RN7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H20 ΙT 1333-74-0P, preparation 7782-44-7P, preparation (manuf. of, by electrolysis of water vapor in planar cell) 1333-74-0 HCA RNHydrogen (8CI, 9CI) (CA INDEX NAME) CN H-HRN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME)

o = o

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CC
     72-10 (Electrochemistry)
     Section cross-reference(s): 49, 61
ST
     electrochem redn water vapor cell; planar cell
     water vapor electroredn; electrolytic cell
     water vapor redn; hydrogen oxygen water
     vapor redn; oxygen hydrogen water vapor
     redn
IT
     Reduction, electrochemical
        (of water vapor in planar cells)
IT
     Electrolytic cells
        (planar, for water vapor redn.)
IT
     7440-02-0, uses and miscellaneous
        (cathodes, for water vapor electroredn. in
        planar cell)
     7732-18-5, vapor
IT
        (electrochem. redn. of, in planar cell)
IT
     1314-23-4, uses and miscellaneous
        (electrolyte, stabilized with yttrium oxide, for water
        vapor electroredn. in planar cells)
IT
     11149-64-7
        (in planar cells for water vapor electrochem.
        redn.)
IT
     1333-74-0P, preparation 7782-44-7P, preparation
        (manuf. of, by electrolysis of water vapor in
        planar cell)
IT
     1314-36-9, uses and miscellaneous
        (zirconia electrolyte stabilized with, for water
        vapor electroredn. in planar cells)
     ANSWER 21 OF 22 HCA COPYRIGHT 2006 ACS on STN
90:46055 Electrolytic gas generator for producing a hydrogen-oxygen
             Inoue, Kiyoshi (Inoue-Japax Research Inc., Japan). Jpn.
     flame.
     Kokai Tokkyo Koho JP 53110989 19780928 Showa, 3 pp.
     (Japanese). CODEN: JKXXAF. APPLICATION: JP 1977-26319 19770310.
AB
     An improved electrolytic gas generator for the H-O torch (used for
     example in a plasma welder) employs a pulsed d.c. in the
     electrolysis. By using a pulsed d.c., the quantity of gas produced
     is increased and the mixing of water vapor with
     the gas involved is minimized.
IT
     1333-74-0P, preparation 7782-44-7P, preparation
        (electroprodn. of, for hydrogen-oxygen welding)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
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H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IC C25B001-04 CC 72-12 (Electrochemistry) ST hydrogen oxygen electroprodn welding torch; plasma welding hydrogen oxygen electroprodn IT Torches Welding (plasma, electroprodn. of hydrogen and oxygen for) 1333-74-0P, preparation 7782-44-7P, preparation IT (electroprodn. of, for hydrogen-oxygen welding) ANSWER 22 OF 22 HCA COPYRIGHT 2006 ACS on STN L64 88:165772 Chemical model for the synthesis of living matter precursors. Simionescu, C. I.; Dumitriu, S.; Popa, V. I.; Bulacovschi, V.; Simionescu, B. (Inst. Polytech., Iasi, Rom.). Revue Roumaine de Chimie, 23(1), 89-102 (French) 1978. CODEN: RRCHAX. ISSN: 0035-3930. A chem. model was established for the synthesis of precursors of AB living matter taking into account the compn. of the primeval atm., the energy sources, and the catalysts. The decompn. of CH4/H2O, CH4/NH3, and CH4/NH3/H2O mixts. in high-frequency plasma led to the formation of aldehydes, alcs., acids, ketones, amines, and nitriles, intermediates in the synthesis of biomonomers. influences of different parameters, catalysts, pH, and energy sources were studied. IT **7732-18-5**, reactions (decompn. of, in primordial atm.) RN 7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME) H20 IT1333-74-0P, biological studies 7782-44-7P, biological studies (formation of, in primordial atm.) RN1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-H

RN

CN

7782-44-7 HCA

Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

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CC'
     6-13 (General Biochemistry)
IT
     74-82-8, reactions
                         7664-41-7, reactions 7732-18-5,
     reactions
        (decompn. of, in primordial atm.)
     50-00-0P, biological studies
                                   64-17-5P, biological studies
IT
     64-18-6P, biological studies
                                   64-19-7P, biological studies
     67-56-1P, biological studies
                                   67-63-0P, biological studies
     67-64-1P, biological studies
                                   74-85-1P, biological studies
     74-86-2P, biological studies
                                   74-89-5P, biological studies
     74-90-8P, biological studies
                                   74-98-6P, biological studies
     75-05-8P, biological studies
                                   75-07-0P, biological studies
     75-65-0P, biological studies
                                   78-92-2P 78-93-3P, biological
              106-97-8P, biological studies
                                              106-99-0P, biological
     studies
     studies
              107-13-1P, biological studies
                                              115-11-7P, biological
              124-38-9P, biological studies
     studies
                                              124-40-3P, biological
     studies
              590-18-1P
                          624-64-6P
                                      630-08-0P, biological studies
     1333-74-0P, biological studies 7782-44-7P,
     biological studies
        (formation of, in primordial atm.)
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## => D L65 1-19 CBIB ABS HITSTR HITIND

- L65 ANSWER 1 OF 19 HCA COPYRIGHT 2006 ACS on STN

  142:25868 Method to manufacture composite polymer electrolyte

  membranes coated with inorganic thin films for fuel cells.

  Ha, Heung-Yong; Kwak, Soon Jong; Kim, Daejin; Shim, Juno; Oh,

  In-Hwan; Hong, Seong-Ahn; Lim, Tae-Hoon; Nam, Suk-Woo (Korea

  Institute of Science and Technology, S. Korea). U.S. Pat. Appl.

  Publ. US 2004241520 A1 20041202, 9 pp. (English). CODEN: USXXCO.

  APPLICATION: US 2003-751138 20031230. PRIORITY: KR 2003-35127

  20030531.
- The present invention relates to a method for manufg. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells using a plasma enhanced CVD (PECVD) method or a reactive sputtering method, so as to reduce the crossover of methanol through polymer electrolyte membranes for fuel cells and enhance the performance of the fuel cells. The manufg. method of composite polymer electrolyte membranes coated with inorg. thin films for fuel cells according to the present invention is characterized to obtain composite membranes by coating the surface of com. composite polymer electrolyte membranes for fuel cells with inorg. thin films using a PECVD method or a reactive sputtering method. The

inorg. materials to form the inorg. thin films are chosen one or more from the group comprising silicon oxide (SiO2), titanium oxide (TiO2), zirconium oxide (ZrO2), zirconium phosphate (Zr(HPO4)2), zeolite, silicalite, and aluminum oxide (Al2O3). The present invention, by coating the polymer electrolyte membranes for fuel cells with inorg. thin films via a PECVD method or a reactive sputtering method, reduces the methanol crossover sizably without seriously reducing the ionic cond. of polymer electrolyte membranes, thereby, when applied to fuel cells, realizes a high performance of fuel cells.

IT 1333-74-0, Hydrogen, uses 7782-44-7, Oxygen, uses (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IC ICM H01M008-10

ICS C08J005-22; H01M004-88; B05D005-12

INCL 429033000; 521027000; 427115000; 502101000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38, 39, 56, 72, 76

ST manuf composite polymer electrolyte membrane coated oxide thin film; reactive sputtering film fuel cell polymer composite electrolyte membrane; plasma enhanced chem vapor deposition film fuel cell membrane

IT Ion exchange membranes

(Aciplex and Dow-type membranes, sulfo-contg.; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Membranes, nonbiological

(composite; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Fluoropolymers, uses

(ethene-propene copolymers; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Polyoxyalkylenes, uses

(fluorine- and sulfo-contg., ionomers, Nafion; method to manuf. composite polymer electrolyte membranes coated with

inorg, thin films for fuel cells) IT Ionomers (fluoropolymers, sulfo-contg., styrene-based, BAM type membranes; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Fluoropolymers, uses (ionomers, sulfo-contg., styrene-based, BAM type membranes; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Polymer electrolytes Reactive sputtering Steam Ultrathin films (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) ITSilicalites (zeolites) Zeolite-group minerals (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Ionic conductivity (of composite polymer electrolyte membranes; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Permeability (of methanol through composite polymer electrolyte membranes; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) ΙT Vapor deposition process (plasma; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Polyketones ' (polyether-, sulfonated; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Polysulfones, uses (polyimide-, sulfonated; method to manuf. composite polymer electrolyte membranes coated with inorq. thin films for fuel cells) IT Polyethers, uses (polyketone-, sulfonated; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells) IT Membranes, nonbiological (polymer electrolyte, Flemion; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

(polymer electrolyte; method to manuf. composite polymer

IT

Fuel cells

electrolyte membranes coated with inorg. thin films for fuel cells)

IT Fluoropolymers, uses

(polyoxyalkylene-, sulfo-contg., ionomers, Nafion; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Ionomers

(polyoxyalkylenes, fluorine- and sulfo-contg., Nafion; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Polyimides, uses

(polysulfone-, sulfonated; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Microwave heating

(sputtering with; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Phenolic resins, uses

(sulfo-contg.; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Polyamides, uses

Polybenzimidazoles

Polymers, uses

(sulfonated; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Oxides (inorganic), uses

(thin films on polymer membrane; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT Membrane electrodes

(using composite polymer electrolyte membranes; method to manuf. composite polymer electrolyte membranes coated with inorq. thin films for fuel cells)

IT 7631-86-9P, Silica, uses

(coating on polymer membrane and targets; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

IT 1344-28-1P, Alumina, uses

(coating on polymer membrane; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

- IT 1333-74-0, Hydrogen, uses 7782-44-7, Oxygen, uses (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- IT 1314-23-4, Zirconium oxide (ZrO2), uses 9002-84-0D, Poly(tetrafluoroethene), sulfonated 9002-88-4D, Polyethylene, sulfonated 9003-07-0D, Polypropylene, sulfonated 9003-53-6D, Polystyrene, sulfonated 13463-67-7, Titanium oxide (TiO2), uses

13772-29-7, Zirconium phosphate (Zr(HPO4)2) 24937-79-9D, Polyvinylidene fluoride, sulfonated 53232-35-2 190673-42-8, Gore-Select

(method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

- IT 7440-37-1, Argon, uses
  - (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- IT 67-56-1, Methanol, uses
  - (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- IT 555-31-7, Aluminum isopropoxide 555-75-9, Aluminum 681-84-5, Tetramethylorthosilicate ethoxide 865-31-6, Aluminum 1450-14-2, Hexamethyldisilane 1992-48-9, methoxide Tetra-isopropylorthosilicate 3085-30-1, Aluminum butoxide 3087-36-3, Titanium ethoxide 4766-57-8, Tetrabutyl orthosilicate 5593-70-4 7245-18-3, Titanium methoxide 23355-24-0, Titanium butoxide

(method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

- Titanium, uses 7440-21-3, Silicon, uses 7440-32-6, Titanium, uses 7440-67-7, Zirconium, uses 14515-04-9 (method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- IT 77950-55-1, Nafion 115
  - (plain and composites with silica; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- IT 78-10-4, Silicon ethoxide 7440-21-3D, Silicon, org. compds.
  7440-32-6D, Titanium, org. compds. 7440-67-7D, Zirconium, org. compds.

(precursor; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)

- IT 7429-90-5D, Aluminum, org. compds.
  - (thin films on polymer membrane; method to manuf. composite polymer electrolyte membranes coated with inorg. thin films for fuel cells)
- L65 ANSWER 2 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 142:24666 Coating and surface treatment processes for strongly adherent surface coatings. Wolf, Jean-Pierre; Kunz, Martin (Ciba Specialty Chemicals Holding Inc., Switz.). PCT Int. Appl. WO 2004103580 Al 20041202, 72 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN,

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YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK,
     ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN,
     TD, TG, TR.
                  (English). CODEN: PIXXD2. APPLICATION: WO
                             PRIORITY: CH 2003-928 20030523.
     2004-EP50806 20040514.
AB
     In a process for the prodn. of a strongly adherent coating on an
     inorg. or org. substrate, wherein (a), a low-temp. plasma
     treatment, a corona discharge treatment or a flame treatment is
     carried out on the inorg. or org. substrate, (b) one or more
     photoinitiators or mixts. of photoinitiators with monomers or/and
     oligomers, contg. at least one ethylenically unsatd. group, or
     solns., suspensions or emulsions of the afore-mentioned substances,
     are applied to the inorg. or org. substrate, and optionally, (c)
     using suitable methods those afore-mentioned substances are dried
     and/or are irradiated with electromagnetic waves
     , it proves advantageous to use compds. of formula (I), (II),
     and/or (IV), IN-L-RG (I), IN-L-RG1-L1-H (II), IN-L-RG1-L1-IN1 (III),
     IN-L-RG1-L1-RG2-L2-IN1 (IV), wherein IN and IN1 are each
     independently of the others a monacylphosphine, monoacylphosphine
     oxide or monoacylphosphine sulfide photoinitiator group; L1 L1 and
     L2 are a single bond or a spacer group; RG is a monovalent radical
     having at least one ethylenically unsatd. C = C bond; and RG1 and
     RG2 are each independently of the other a divalent radical having at
     least one ethylenically unsatd. C = C bond.
IT
     1333-74-0, Hydrogen, uses 7732-18-5, Water
     , uses 7782-44-7, Oxygen, uses
        (plasma gas; coating and surface treatment
        processes for strongly adherent surface coatings)
RN
     1333-74-0 HCA
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-- H
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H<sub>2</sub>O
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
IC
     ICM
         B05D003-14
```

ICS B05D003-08; C07F009-50; C07F009-53 42-2 (Coatings, Inks, and Related Products)

CC

IT Coating materials
Coating process
Electric corona
Flame

## Plasma

(coating and surface treatment processes for strongly adherent surface coatings)

IT Air

(plasma gas; coating and surface treatment processes for strongly adherent surface coatings)

IT 124-38-9, Carbon dioxide, uses 1333-74-0, Hydrogen, uses
7439-90-9, Krypton, uses 7440-37-1, Argon, uses 7440-59-7,
Helium, uses 7440-63-3, Xenon, uses 7727-37-9, Nitrogen, uses
7732-18-5, Water, uses 7782-44-7,
Oxygen, uses

(plasma gas; coating and surface treatment processes for strongly adherent surface coatings)

L65 ANSWER 3 OF 19 HCA COPYRIGHT 2006 ACS on STN
140:345634 Modeling of chemical kinetics in inductively coupled
plasma torches. Pelletier, D.; Delannoy, Y.; Proulx, P.
(EPM, ENSHMG, Saint Martin d'Heres, F-38402, Fr.). International
Conference on Electromagnetic Processing of Materials, 4th, Lyon,
France, Oct. 14-17, 2003, 330-335. Forum Editions: Paris, Fr.
(English) 2003. CODEN: 69FIFN.

This paper presents some of the preliminary numerical results obtained from a two-dimensional FLUENT based model to calc. the electromagnetic field and the reactive flow in inductively coupled plasma torches. Six different species (H2, O2, H2O, OH, H, O) were introduced together with four reversible chem. reactions to represent a transferred plasma used for silicon purifn. The model shows that a significant quantity of reactive radicals (OH, O, H) is present near the liq. silicon target, increasing in this way the probability to produce volatile compds. at the free surface by interaction with dissolved impurities, and so to speed up the purifn. process.

IT 1333-74-0, Hydrogen, reactions 7732-18-5, Water, reactions 7782-44-7, Oxygen, reactions (modeling of chem. kinetics in inductively coupled plasma torches)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME)

```
H<sub>2</sub>O
```

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

o = 0

CC 67-3 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms)

Section cross-reference(s): 65, 76

- ST modeling chem kinetics inductively coupled **plasma** torch silicon purifn
- IT Plasma torches

(inductively coupled; modeling of chem. kinetics in inductively coupled plasma torches)

IT Plasma

Reaction kinetics
Simulation and Modeling
Surface reaction

(modeling of chem. kinetics in inductively coupled plasma torches)

IT Radicals, reactions

(modeling of chem. kinetics in inductively coupled plasma torches)

- 1T 1333-74-0, Hydrogen, reactions 3352-57-6, Hydroxyl,
   reactions 7732-18-5, Water, reactions 7782-44-7,
   Oxygen, reactions 12385-13-6, Atomic hydrogen, reactions
   17778-80-2, Atomic oxygen, reactions
   (modeling of chem. kinetics in inductively coupled plasma torches)
- TT 7440-21-3P, Silicon, processes (modeling of chem. kinetics in inductively coupled plasma torches)
- L65 ANSWER 4 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 139:335066 Method and apparatus for plasma deposition of chemically reactive groups on substrates chemically reactive substrates obtainable by the method and use thereof. Christensen, Soren Flygenring; Petersen, Steen Guldager (NKT Research & Innovation A/s, Den.). PCT Int. Appl. WO 2003090939 A1

  20031106, 70 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG,

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SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
     ZW; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR,
     GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR.
     (English).
                CODEN: PIXXD2. APPLICATION: WO 2003-DK272 20030425.
     PRIORITY: DK 2002-637 20020425.
AΒ
     The present invention relates to a method and app. for
     plasma deposition of a chem. reactive group (Y-Z) on a
     substrate, chem. reactive substrates, and use thereof, e.g. for
     immobilization of biomols.; the method comprising: (a) providing at
     least one precursor (A-X (Y)) for the chem. reactive group; (b)
     providing at least one donor (D(Z)), said at least one donor
     comprising at least one addn. group (Z), optionally said at least
     one addn. group (Z) being comprised in said precursor (A-X (Y)) and
     optionally said at least one donor (D(Z)) is not being provided; (c)
     providing a substrate (M); (d) providing a gas plasma,
     said gas plasma having a pressure and an energy to form at
     least one activated carrier group (B); and (e) reacting said
     substrate (M), said at least one precursor (A-X (Y)), said at least
     one donor (D(Z)) in said gas plasma so that said chem.
     reactive group (Y-Z) is bound to said substrate, either directly
     (M-Y-Z) or via said at least one activated carrier group (M-B-Y-Z),
     and so that when exposed to a substance which chem. reacts with said
     chem. reactive group, said substance binds thereto.
ΙT
     1333-74-0, Hydrogen, reactions 7732-18-5, Water,
     reactions 7782-44-7D, Oxygen, compds. contg.
        (method and app. for plasma deposition of chem.
        reactive groups on substrates chem. reactive substrates
        obtainable by the method and use thereof)
RN
     1333-74-0 HCA
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-H
RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
o = o
```

IC

ICM B05D007-24

ICS A61L033-00; H05H001-24; H01J037-32

CC 9-1 (Biochemical Methods)

app plasma deposition chem reactive group substrate

IT Functional groups

(Active; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Frequency

ST

(Audio; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Functional groups

Molecules

(Chem. reactive; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Aldehydes, reactions

Esters, reactions

Ketones, reactions

(Chlorinated; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Functional groups

(Cleaving; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Sensors

(DNA; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Molecules

(Environmental; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Frequency

(Microwave; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Thioethers

(Satd. heterocyclic; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Nitriles, reactions

(Satd.; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Thioethers

(Unsatd. heterocyclic; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive

substrates obtainable by the method and use thereof) IT Thioethers (Unsatd. substituted heterocyclic; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof) Nitriles, reactions IT Thioethers (Unsatd.; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof) Nitriles, reactions IT (arom.; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof) IT **Apparatus** Atoms Bond Bond cleavage Cantilevers (components) Carbonyl group Carriers Containers Crystals Electric current Electric insulators Electrodes Energy Frequency Gases Holders Immobilization, molecular or cellular Membranes, nonbiological Pipes and Tubes Plasma Plates Pressure Reaction Sensors Spheres Sulfhydryl group Vacuum Vacuum pumps Wires (method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT

DNA

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Biochemical compounds

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Carboxylic acids, reactions

(method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Esters, reactions

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Fibers

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Lactones

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Monomers

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Nitriles, reactions

(method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Electric generators

(power supplies; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary, Heterocyclic, satd.; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary, Satd. substituted heterocyclic; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use

thereof)

IT Amines, reactions

(secondary, Satd.; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary, Unsatd. heterocyclic; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary, Unsatd. substituted heterocyclic; method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary, Unsatd.; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(secondary; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Amines, reactions

(tertiary; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT Esters, reactions

(unsatd.; method and app. for **plasma** deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

IT 74-82-8, Methane, reactions 75-00-3, Ethyl chloride Acetonitrile, reactions 75-43-4, Dichlorofluoromethane 75-69-4, Trichlorofluoromethane Carbonyl chloride 80-62-6. 96-54-8, 1-Methylpyrrole Methyl methacrylate 97-62-1, Ethyl 100-47-0, Benzonitrile, reactions isobutyrate 102-70-5, 107-13-1, Acrylonitrile, reactions Triallylamine 107-47-1, 109-74-0, tert-Butyl sulfide 108-29-2,  $\gamma$ -Valerolactone n-Butanenitrile 109-89-7, Diethylamine, reactions 109-97-7, Pyrrole 110-01-0, Tetrahydrothiophene 110-02-1, Thiophene 110-86-1, Pyridine, reactions 110-89-4, Piperidine, reactions 120-94-5, 1-Methylpyrrolidine 121-44-8, Triethylamine, reactions 123-75-1, Pyrrolidine, reactions 124-02-7, Diallylamine 141-78-6, Ethyl acetate, reactions 288-13-1, Pyrazole 288-32-4, Imidazole, reactions 289-95-2, Pyrimidine 547-63-7, Methyl 554-14-3, 2-Methylthiophene isobutyrate 592-88-1, Allyl sulfide 616-43-3, 3-Methylpyrrole 623-47-2, Ethyl propiolate 2,4-Dimethylpyrrole 627-37-2, N-Allylmethylamine 638-02-8,

2,5-Dimethylthiophene 922-67-8, Methyl propiolate 1072-63-5, 1300-21-6, Dichloroethane 1333-74-0, N-Vinylimidazole Hydrogen, reactions 3068-88-0, β-Butyrolactone 7664-41-7, 7704-34-9D, Sulfur, compds. contg. Ammonia, reactions 7727-37-9D, Nitrogen, compds. contq. 7732-18-5, Water, reactions 7782-44-7D, Oxygen, compds. contg. 7782-50-5D, 10152-76-8, Allyl methyl sulfide Chlorine, mols. contg. 26638-19-7, Dichloropropane 26446-76-4, Chloropropane (method and app. for plasma deposition of chem. reactive groups on substrates chem. reactive substrates obtainable by the method and use thereof)

L65 ANSWER 5 OF 19 HCA COPYRIGHT 2006 ACS on STN
139:8735 nanopowder production system. Wu, L. W.; Huang, Wen-Chiang
(USA). U.S. Pat. Appl. Publ. US 2003108459 A1 20030612,
14 pp. (English). CODEN: USXXCO. APPLICATION: US 2001-6378
20011210.

AB A system for synthesizing a nano-scaled powder material, includes several sub-systems: (A) a chamber for contg. nano-scaled cluster generating devices from a material selected from the group consisting of a metal, a metal compd., and a ceramic; (B) a twin-wire electrode device disposed within this chamber with this electrode device including: (1) two wires made up of this material, each having a leading tip and each being continuously or intermittently fed into the chamber in such a fashion that the two leading tips are maintained at a desired sepn.; and (2) power supply for providing elec. current and gas supply for providing a working gas flow for creating an ionized arc between the two leading tips for melting and/or vaporizing the material to generate nano-scaled clusters; (C) devices for injecting a quench gas and/or a reaction gas into a quenching/reaction zone inside the chamber at a point downstream from the arc to produce nano-scaled powder particles; and (D) devices such as a cyclone and powder classifier to collect the nano-scaled powder material.

IT 7553-56-2, Iodine, processes 7782-44-7, Oxygen, processes

(reaction gas; nanopowder prodn. system)

RN 7553-56-2 HCA

CN Iodine (8CI, 9CI) (CA INDEX NAME)

I-I

RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME)

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IT
     1333-74-0, Hydrogen, uses
        (working gas; nanopowder prodn. system)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI)
CN
                          (CA INDEX NAME)
H-H
IC
     ICM B01J019-08
INCL 422186040; 422186290
     48-11 (Unit Operations and Processes)
CC
IT
     Plasma
        (arc; nanopowder prodn. system)
     Air
IT
       Water vapor
        (quench gas; nanopowder prodn. system)
IT
     7440-38-2, Arsenic, processes 7553-56-2, Iodine, processes
     7704-34-9, Sulfur, processes
                                    7723-14-0, Phosphorus, processes
     7726-95-6, Bromine, processes 7727-37-9, Nitrogen, processes
     7782-41-4, Fluorine, processes 7782-44-7, Oxygen,
     processes 7782-49-2, Selenium, processes
                                                  7782-50-5, Chlorine,
                 13494-80-9, Tellurium, processes
    processes
        (reaction gas; nanopowder prodn. system)
     1333-74-0, Hydrogen, uses
        (working gas; nanopowder prodn. system)
    ANSWER 6 OF 19 HCA COPYRIGHT 2006 ACS on STN
138:361446 Design of field emission devices using modified carbon
     nanotubes. Takai, Mikio; Fischer, Alan B.; Niu, Chunming; Tennent,
     Howard G.; Hoch, Robert; Biebuyck, Hans (Hyperion Catalysis
     International, Inc., USA). U.S. Pat. Appl. Publ. US 2003090190 A1
     20030515, 33 pp.
                       (English). CODEN: USXXCO. APPLICATION:
     US 2002-171760 20020614. PRIORITY: US 2001-298193P 20010614.
     The invention relates to the design of field emission devices using
AB
    modified carbon nanotubes, where the carbon nanotubes are used to
     form the cathode.
                        The device consists of an anode and a cathode,
     where the cathode includes carbon nanotubes, which have been
     subjected to an energy, plasma, chem., or mech. treatment.
IT
     7782-44-7, Oxygen, processes
        (ions, plasma carbon nanotubes treated with; design of
        field emission devices using modified carbon nanotubes)
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
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IT 1333-74-0, Hydrogen, processes (ions, plasma, carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes) 1333-74-0 HCA RNHydrogen (8CI, 9CI) (CA INDEX NAME) CN H-HIT **7732-18-5**, Water, processes (plasma, carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes) RN 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H<sub>2</sub>O IC ICM H01J001-05 ICS H01J001-38 INCL 313311000 76-12 (Electric Phenomena) Section cross-reference(s): 41, 73, 74 ΙT Atomic beams Chemisorbed substances Chemisorption Electromagnetic wave Electron beams Electrophiles Gamma ray Heat treatment Ion beams Ionizing radiation Laser radiation Microwave Molecular beams Nucleophiles Oxidizing agents Plasma Reducing agents UV radiation X-ray (carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes) IT 7782-44-7, Oxygen, processes (ions, plasma carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes) IT 1333-74-0, Hydrogen, processes 7440-37-1, Argon, processes

7440-59-7, Helium, processes (ions, plasma, carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes) 74-85-1, Ethylene, processes IT 75-46-7, Fluoroform Bromo-trifluoromethane 75-71-8, Difluoro-dichloromethane 75-72-9, Chlorotrifluoromethane 75-73-0, Carbon tetrafluoride 116-14-3, Perfluoroethylene, processes 2551-62-4, Sulfur 7664-41-7, Ammonia, processes hexafluoride 7727-37-9, Nitrogen, processes 7732-18-5, Water, processes (plasma, carbon nanotubes treated with; design of field emission devices using modified carbon nanotubes)

L65 ANSWER 7 OF 19 HCA COPYRIGHT 2006 ACS on STN 137:248831 Production of metal films on polymer surfaces treated with cold plasma to reduce contact angle. Ryu, Seung-Kyun; Su, Kon (S. Korea). PCT Int. Appl. WO 2002075020 A1 20020926, 32 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, (English). CODEN: PIXXD2. APPLICATION: WO 2002-KR382 TD, TG, TR. PRIORITY: KR 2001-11334 20010306. 20020306. AB A method for plating the surface of a polymer material with a metal film comprises treatment of the polymer surface with cold plasma to introduce hydrophilic functional groups and plating the surface of the polymer material with a metal film according to the electroless plating method. The cold plasma treatment reduces water contact angle of the polymer surface by 5 to 60°. Addnl. step of immersing the polymer material into an org. solvent for 0.1 to 5 min or of washing the polymer material with ultrasonic washing machine can be used after introducing the hydrophilic functional groups by cold plasma The method is very stable and can endow new properties -treatment. to the polymer material, such as improvement of appearance, mech. properties, heat resistance, rendering heat and elec. cond., soldering ability and electromagnetic wave shielding effect. The polymer materials plated by this method can be used for shielding electromagnetic waves generated from the information processing equipment, prepg. printed circuit boards, electrodes and antistatic boards. Thus, the surface of polypropylene (PP) was treated for 3 min at 15° with cold plasma generated in the mixt. of acrylic acid, oxygen and methane. The water contact angle of PP was reduced from 90° to 8.5°. The treated polymer was used to form copper film on

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the surface by electroless plating using palladium as catalytically
     activated phase.
IT
     1333-74-0, Hydrogen, uses 7732-18-5, Water, uses
     7782-44-7, Oxygen, uses
        (cold plasma; prodn. of metal films on polymer surfaces
        treated with cold plasma to reduce contact angle)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H20
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
o = o
IC
     ICM C23C016-44
     38-3 (Plastics Fabrication and Uses)
CC
     Section cross-reference(s): 76
     electroless plating metal coating treated polymer surface; polymer
ST
     cold plasma treatment hydrophilic surface metal film prodn
IT
     Air
        (cold plasma; prodn. of metal films on polymer surfaces
        treated with cold plasma to reduce contact angle)
     Coating process
IT
        (electroless; prodn. of metal films on polymer surfaces treated
        with cold plasma to reduce contact angle)
     Printed circuit boards
IT
        (metal films on cold plasma-treated polymer surfaces
        for prodn. of)
IT
     Cold plasma
        (prodn. of metal films on polymer surfaces treated with cold
       plasma to reduce contact angle)
IT
     Polycarbonates, uses
     Polyesters, uses
     Polyimides, uses
     Silicone rubber, uses
        (prodn. of metal films on polymer surfaces treated with cold
       plasma to reduce contact angle)
     Acrylic polymers, uses
IT
```

Epoxy resins, uses Fluoropolymers, uses Rubber, uses

(prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)

IT 50-00-0, Formalin, uses 67-64-1, Acetone, uses 74-82-8, Methane,
 uses 75-07-0, Acetaldehyde, uses 79-10-7, Acrylic acid, uses
 1333-74-0, Hydrogen, uses 7440-37-1, Argon, uses
 7727-37-9, Nitrogen, uses 7732-18-5, Water, uses
 7782-44-7, Oxygen, uses

(cold plasma; prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)

IT 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-31-5, Tin, uses

(in prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)

- IT 9002-84-0, Polytetrafluoroethylene 9002-86-2, Polyvinyl chloride 9002-88-4, Polyethylene 9003-07-0, Polypropylene 9003-53-6, Polystyrene 25038-59-9, Poly(ethylene terephthalate), uses (prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)
- IT 9003-56-9, Acrylonitrile-butadiene-styrene copolymer (prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)
- IT 7440-02-0, Nickel, uses 7440-48-4, Cobalt, uses 7440-50-8, Copper, uses

(prodn. of metal films on polymer surfaces treated with cold plasma to reduce contact angle)

L65 ANSWER 8 OF 19 HCA COPYRIGHT 2006 ACS on STN

- 136:202788 Conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge **plasma** method. Okumoto, Mamoru; Mizuno, Akira (Department of Ecological Engineering, Toyohashi University of Technology, Toyohashi, Tempaku-cho, Aichi, 441-8580, Japan). Catalysis Today, 71(1-2), 211-217 (English) 2001. CODEN: CATTEA. ISSN: 0920-5861. Publisher: Elsevier Science B.V..
- AB This paper presents new conversion method of methane to higher hydrocarbon fuels such as MeOH (methanol), formaldehyde, using pulsed discharge plasma under room temp. and atm. pressure. The expts. were carried out with special attention to the effect of the specific input energy (SIE) defined as the elec. input energy per unit mass of the material gas. In the study of partial oxidn. of methane, exptl. results indicated that the methanol and formaldehyde prodn. has an optimum SIE value. The highest methanol and formaldehyde prodn. ability and reaction selectivity were achieved with relatively low SIE of 360 J/l, based on feed gas. Under this optimum condition, a max. prodn. ability of .apprx.0.65 μmol/J and selectivity of 64% were obtained. However, to achieve

IT

RNCN

H-H

RN

CN

H<sub>2</sub>O

IT

RN CN

I-I

RN

CN

CC

ST

IT

IT

IT

selective reaction of methane, conversion of methane with halogen materials to Me halide such as Me iodine was also studied. expt., max. Me iodine prodn. selectivity of 95% was achieved with the prodn. ability of 0.45  $\mu$ mol/J. 1333-74-0, Hydrogen, formation (nonpreparative) 7732-18-5, Water, formation (nonpreparative) (conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 7553-56-2, Iodine, reactions 7782-44-7, Oxygen, reactions (conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method) 7553-56-2 HCA Iodine (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 051-11 (Fossil Fuels, Derivatives, and Related Products) Section cross-reference(s): 45 methane hydrocarbon fuel pulsed discharge plasma partial oxidn; methanol formaldehyde methyl iodide (conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method) Hydrocarbons, formation (nonpreparative) (conversion of methane for higher hydrocarbon fuel synthesis

(conversion of methane for higher hydrocarbon fuel synthesis

using pulsed discharge plasma method)

Natural gas, reactions

using pulsed discharge plasma method)

IT Halogenation

(for CH4 oxidn. selectivity improvement; conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)

IT Hydrocarbons, preparation

(oxy; conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)

IT Oxidation

(partial; conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge **plasma** method)

IT Reactors

(plasma; conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)

IT Electric discharge

(pulsed; conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)

TT 74-84-0, Ethane, formation (nonpreparative) 74-85-1, Ethylene,
 formation (nonpreparative) 75-07-0, Acetaldehyde, formation
 (nonpreparative) 124-38-9, Carbon dioxide, formation
 (nonpreparative) 630-08-0, Carbon monoxide, formation
 (nonpreparative) 1333-74-0, Hydrogen, formation
 (nonpreparative) 7732-18-5, Water, formation
 (nonpreparative)

(conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)

- IT 50-00-0P, Formaldehyde, preparation 67-56-1P, Methanol, preparation 74-88-4P, Methyl iodide, preparation (conversion of methane for higher hydrocarbon fuel synthesis using pulsed discharge plasma method)
- L65 ANSWER 9 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 136:143637 Method of forming a metal wiring in a semiconductor device. Pyo, Sung Gyu (S. Korea). U.S. Pat. Appl. Publ. US 2002009884 A1 20020124, 6 pp. (English). CODEN: USXXCO. APPLICATION: US 2001-874505 20010605. PRIORITY: KR 2000-33984 20000620.
- AB A method of forming a metal wiring in a semiconductor device is presented. A Cu wiring is formed by means of CECVD method by which a chem. enhancer layer is used for increasing the deposition speed of Cu. The damascene pattern is filled by means of MOCVD method using a Cu precursor to increase the deposition speed. The chem. enhancer layer rises to the surface of Cu after deposition of Cu by a CECVD method and then the relatively high resistivity chem. enhancer layer that has risen to the surface of Cu by plasma

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method)

process is removed. Therefore, the ultra-fine damascene pattern can be rapidly filled with Cu without increasing the resistance of the Cu wiring. **7553-56-2**, Iodine, uses (catalyst for chem. enhancer layer formation; method of forming a copper metal wiring in a semiconductor device by vapor deposition method) RN 7553-56-2 HCA Iodine (8CI, 9CI) (CA INDEX NAME) CN I-I1333-74-0, Hydrogen, processes 7782-44-7, Oxygen, processes (vapor deposition ambient; method of forming a copper metal wiring in a semiconductor device by vapor deposition method) RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0ICM H01L021-44 INCL 438687000 76-3 (Electric Phenomena) Section cross-reference(s): 29, 75 Water vapor (catalyst for chem. enhancer layer formation; method of forming a copper metal wiring in a semiconductor device by vapor deposition method) Vapor deposition process (plasma; method of forming a copper metal wiring in a semiconductor device by vapor deposition method) 74-88-4, Iodomethane, uses 75-03-6, Iodoethane Diiodomethane 754-05-2, Trimethylvinylsilane 865-50-9 7440-68-8, Astatine, uses 7553-56-2, Iodine, 1522-22-1 7726-95-6, Bromine, uses 7782-41-4, Fluorine, uses 7782-50-5, Chlorine, uses (catalyst for chem. enhancer layer formation; method of forming a

copper metal wiring in a semiconductor device by vapor deposition

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IT
     1333-74-0, Hydrogen, processes 7440-37-1, Argon, processes
     7664-41-7, Ammonia, processes 7727-37-9, Nitrogen, processes
     7782-44-7, Oxygen, processes 10028-15-6, Ozone, processes
        (vapor deposition ambient; method of forming a copper metal
        wiring in a semiconductor device by vapor deposition method)
     ANSWER 10 OF 19 HCA COPYRIGHT 2006 ACS on STN
135:335394 New chemical species of a magnecule. Santilli, Ruggero Maria
     (Hadronic Press, Inc., USA). U.S. Pat. Appl. Publ. US 20010038087
     Al 20011108, 60 pp., Cont.-in-part of U.S. Ser. No.
     586,926. (English). CODEN: USXXCO. APPLICATION: US 2001-826183
     20010404. PRIORITY: US 1994-254377 19940606; US 1997-785797
     19970121; US 1998-106170 19980629; US 1998-133348 19980813; US
     1999-372278 19990811; US 2000-586926 20000605.
     A novel chem. species, called magnecules, which is composed of
AB
     clusters of mols., and/or dimers, and/or atoms formed by internal
     bonds due to the magnetic polarization of the orbits of at least
     some of the peripheral at. electrons present in the cluster, the
     intrinsic magnetic field of nuclei present in the cluster, and the
     intrinsic magnetic fields of valence electrons present in the
     cluster that are not correlated in singlet couplings to other
     electrons to form valence bonds is disclosed.
     1333-74-0, Hydrogen, properties 7732-18-5, Water,
IT
     properties 7782-44-7, Oxygen, properties
        (new chem. species of magnecule composed of clusters of)
RN
     1333-74-0 HCA
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-H
RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
0 = 0
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IT

ICM C09K003-00

Electric furnaces

65-5 (General Physical Chemistry)

Section cross-reference(s): 51, 73, 77

INCL 252062510R

(arc, plasma; new chem. species of magnecule composed of clusters of mols., dimers, and/or atoms formed by passing through)

## IT Electromagnetic field

(new chem. species of magnecule composed of clusters of mols., dimers, and/or atoms with internal bonds due to magnetic polarization by)

- L65 ANSWER 11 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 131:326978 Method for detoxification of moist wastes containing oxidizable organic materials by drying and oxidation under UV irradiation. Laflamme, Claude B.; Labrecque, Raynald (Hydro-Quebec, Can.). PCT Int. Appl. WO 9958459 Al 19991118, 26 pp. DESIGNATED STATES: W: AL, AU, BA, BB, BG, BR, CN, CU, CZ, EE, GD, GE, HR, HU, ID, IL, IN, IS, JP, KP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (French). CODEN: PIXXD2. APPLICATION: WO 1999-CA79 19990129. PRIORITY: CA 1998-2237414 19980511.
- AB Moist org. wastes are oxidized and dried by feeding into a rotary kiln provided with a refractory coating and contg. a heat transferring material; then heating the waste mixed with the heat transferring material at ≥300°C (300-900°C) in the presence of chem. active species (e.g., H3PO4, FeSO4, oxides, hydroxides, carbonates, phosphates) and elec. generated UV irradn. for dewatering and oxidn. of the org. materials with gas formation. The kiln is rotated and heated by a plasma torch, elec. arc, or elec. discharge device. The oxidn. is promoted and catalyzed by the chem. active species and the UV radiation. The gases are evacuated from the kiln. The method provides inexpensive destruction of polluting and/or oxidizable toxic materials.
- IT 7664-38-2, Phosphoric acid, uses

(drying and oxidn. of moist org. wastes in rotary kilns under UV irradn.)

- RN 7664-38-2 HCA
- CN Phosphoric acid (7CI, 8CI, 9CI) (CA INDEX NAME)

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HO- b- OH
   OH
IT
     1333-74-0, Hydrogen, uses 7727-37-9, Nitrogen,
     uses 7782-44-7, Oxygen, uses
        (plasma torch; drying and oxidn. of moist org. wastes
        in rotary kilns under UV irradn.)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7727-37-9 HCA
CN
     Nitrogen (8CI, 9CI)
                           (CA INDEX NAME)
     7782-44-7 HCA
RN
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
IT
     7732-18-5, Water, uses
        (vapor, plasma torch; drying and oxidn. of
        moist org. wastes in rotary kilns under UV irradn.)
     7732-18-5 HCA
RN
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
IC
     ICM
          C02F011-06
     ICS C02F001-32; F23G005-10; F23G005-20; F23G007-00; A62D003-00
CC
     60-4 (Waste Treatment and Disposal)
     7664-38-2, Phosphoric acid, uses
IT
     7720-78-7, Ferrous sulfate
        (drying and oxidn. of moist org. wastes in rotary kilns under UV
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irradn.)

IT 124-38-9, Carbon dioxide, uses 74-82-8, Methane, uses 630-08-0, Carbon monoxide, uses 1333-74-0, Hydrogen, uses 7440-37-1, Argon, uses **7727-37-9**, Nitrogen, uses 7782-44-7, Oxygen, uses (plasma torch; drying and oxidn. of moist org. wastes in rotary kilns under UV irradn.) IT **7732-18-5**, **Water**, uses (vapor, plasma torch; drying and oxidn. of moist org. wastes in rotary kilns under UV irradn.) ANSWER 12 OF 19 HCA COPYRIGHT 2006 ACS on STN 131:280224 Etching low-k dielectrics. Yan, Chun; Hsueh, Gary C.; Ye, Yan; Ma, Diana Xiaobing (Applied Materials, Inc., USA). PCT Int. Appl. WO 9952135 A1 19991014, 45 pp. DESIGNATED STATES: W: JP, KR, SG; RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1999-US7152 19990331. PRIORITY: US 1998-54285 19980402. The disclosure pertains to a method for plasma etching of AB low-k materials, particularly org. polymer-based low-k materials. The method employs an etchant plasma where the major etchant species are generated from a halogen other than F and O. The preferred halogen is Cl. The volumetric (flow rate) or at. ratio of the halogen: O in the plasma source gas ranges from .apprx.1:20 to .apprx.20:1. When the halogen is Cl, the preferred at. ratio of Cl:O ranges from .apprx.1:10 to .apprx.5:1. When this at. ratio for Cl:O is used, the etch selectivity for the low-k material over adjacent O-comprising or N-comprising layers is advantageous, typically .gtorsim.10:1. The plasma source gas may contain additives in an amt. of ≤15% by vol. which are designed to improve selectivity for the low-k dielec. over an adjacent material, to provide a better etch profile, or to provide better crit. dimension control. IT 1333-74-0, Hydrogen, processes 7553-56-2, Iodine, processes 7732-18-5, Water, processes 7782-44-7, Oxygen, processes (plasma etching low-k dielecs. in gases contg.) RN1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-H7553-56-2 HCA RN

CN

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Iodine (8CI, 9CI) (CA INDEX NAME)

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RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI)
                        (CA INDEX NAME)
H20
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
IC
     ICM H01L021-311
CC
     76-3 (Electric Phenomena)
     Section cross-reference(s): 38
ST
     plasma etching dielec; halogen oxygen plasma
     etching polymeric dielec
IT
     Polyethers, processes
        (arom., fluorinated; plasma etching low-k dielecs.
        from)
IT
     Polyethers, processes
        (arom., oxazoles; plasma etching low-k dielecs. from)
ΙT
     Polyethers, processes
        (arom., poly(arylene) ethers; plasma etching low-k
        dielecs. from)
     Electric insulators
IT
        (plasma etching low-k dielecs.)
IT
     Fluoropolymers, processes
     Polybenzoxazoles
     Polyimides, processes
     Polymers, processes
     Polyoxyphenylenes
        (plasma etching low-k dielecs. from)
     Halogens
IT
     Hydrocarbons, processes
        (plasma etching low-k dielecs. in gases contg.)
IT
     Etching
        (plasma; plasma etching low-k dielecs.)
IT
     Polyguinoxalines
        (polyphenylquinoxalines; plasma etching low-k dielecs.
        from)
IT
     Etching
        (selective; in plasma etching low-k dielecs.)
IT
     25669-37-8
        (Parylene AF, Parylene F; plasma etching low-k dielecs.
     697-11-0, Perfluorocyclobutene
IT
                                      3345-29-7
                                                   9002-84-0
                                                               9002-88-4,
     Polyethylene 9003-07-0, Polypropylene 9003-53-6, Polystyrene
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25722-33-2, Parylene-N

25038-76-0, Polynorbornene

low-k dielecs. from)

36605-57-9

124221-30-3, Cyclotene 181997-50-2, FPI-136M 203945-07-7, SiLK 213329-13-6, Flare 2.0 (plasma etching low-k dielecs. from) IT 56-23-5, processes 64-17-5, Ethanol, processes 67-56-1, Methanol, processes 67-66-3, Chloroform, processes Bromomethane, processes 75-44-5, Phosgene 75-46-7, Trifluoromethane 75-73-0 76-06-2, Trichloronitromethane 124-38-9, Carbon dioxide, processes 630-08-0, Carbon monoxide, 1300-21-6, Dichloroethane 1333-74-0, Hydrogen, 7439-90-9, Krypton, processes 7440-01-9, Neon, processes 7440-37-1, Argon, processes processes 7440-59-7, Helium, 7440-63-3, Xenon, processes **7553-56-2**, Iodine, processes 7647-01-0, Hydrogen chloride, processes processes 7727-37-9, Nitrogen, processes Bromine, processes 7732-18-5, Water, processes 7782-44-7, Oxygen, processes 7782-50-5, Chlorine, processes 7783-54-2, Nitrogen 7789-33-5, Iodine bromide (IBr) 7790-99-0, Iodine fluoride (NF3) chloride (ICl) 10024-97-2, Nitrogen oxide (N2O), processes 10026-04-7, Silicon chloride (SiCl4) 10028-15-6, Ozone, processes 10034-85-2, Hydrogen iodide 10035-10-6, Hydrogen bromide, 10102-44-0, Nitrogen dioxide, processes 10294-34-5, processes Boron chloride (BCl3) 13863-41-7, Bromine chloride (BrCl) 25323-30-2, Dichloroethene 25323-89-1, Trichloroethane 55299-13-3, Dibromodichloroethane (plasma etching low-k dielecs. in gases contg.) 7631-86-9, Silica, processes 11105-01-4, Silicon nitride oxide IT 12033-89-5, Silicon nitride, processes (plasma etching low-k dielecs. in relation to) IT 25135-16-4D, Polynaphthalene, derivs.

L65 ANSWER 13 OF 19 HCA COPYRIGHT 2006 ACS on STN

130:260787 Cleaning of contamination from electron-emissive elements.

Knall, N. Johan; Porter, John D.; Stanners, Colin D.; Spindt,
Christopher J.; Bascom, Victoria A. (Candescent Technologies
Corporation, USA). PCT Int. Appl. WO 9917323 A2 19990408,
38 pp. DESIGNATED STATES: W: JP, KR; RW: AT, BE, CH, CY, DE, DK,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN:
PIXXD2. APPLICATION: WO 1998-US18509 19980922. PRIORITY: US
1997-940873 19970930.

(polynaphthalene-N, polynaphthalene-F; plasma etching

AB Multiple procedures are presented for removing contaminant material from electron-emissive elements of an electron-emitting device. One procedure involves converting the contaminant material into gaseous products, typically by operating the electron-emissive elements, that move away from the electron-emissive elements. Another procedure entails converting the contaminant material into further

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H20

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IT

material and removing the further material. An addnl. procedure involves forming surface coatings over the electron-emissive elements. The contaminant material is then removed directly from the surface coatings or by removing at least part of each surface coating. 1333-74-0, Hydrogen, uses 7553-56-2, Iodine, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses (in cleaning of contamination from electron-emissive elements) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) H-H7553-56-2 HCA Iodine (8CI, 9CI) (CA INDEX NAME) I-I7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) o = 0ICM H01J 76-12 (Electric Phenomena) Section cross-reference(s): 66 (plasma; cleaning of contamination from electron-emissive elements) 50-00-0, Methanal, uses 56-23-5, Carbon tetrachloride, uses 64-18-6, Formic acid, uses 64-19-7, Acetic acid, uses Hexanal 67-56-1, Methanol, uses 67-64-1, Acetone, uses 67-66-3, Trichloromethane, uses 74-82-8, Methane, uses 74-85-1, Ethene, uses 74-86-2, Ethyne, uses 74-87-3, Chloromethane, uses 75-09-2, Dichloromethane, uses 75-10-5, Difluoromethane 75-73-0, Carbon tetrafluoride 79-09-4, Propionic acid, uses 111-27-3, Hexanol, uses 111-65-9, Octane, uses 124-38-9, Carbon dioxide,

302-01-2, Hydrazine, uses 593-53-3, Fluoromethane

630-08-0, Carbon monoxide, uses 1333-74-0, Hydrogen, uses

7439-90-9, Krypton, uses 7440-01-9, Neon, uses 7440-37-1, Argon, 7440-59-7, Helium, uses 7440-63-3, Xenon, uses Sulfur dioxide, uses 7553-56-2, Iodine, uses 7637-07-2, Boron trifluoride, uses 7647-01-0, Hydrogen chloride, uses 7664-39-3, Hydrogen fluoride, uses 7664-41-7, Ammonia, uses 7722-84-1, Hydrogen peroxide, uses 7726-95-6, Bromine, uses 7727-37-9, Nitrogen, uses **7732-18-5**, Water, uses 7782-41-4, Fluorine, uses 7782-44-7, Oxygen, uses 7782-50-5, Chlorine, uses 7783-06-4, Hydrogen sulfide, uses 7783-07-5, Hydrogen selenide 7783-09-7, Hydrogen telluride 7783-54-2, Nitrogen trifluoride 7784-42-1, Arsine 7803-51-2, Phosphorus trihydride 7803-52-3, Stibine 10024-97-2, Nitrous 10028-15-6, Ozone, uses 10034-85-2, Hydrogen iodide oxide, uses 10035-10-6, Hydrogen bromide, uses 10102-43-9, Nitric oxide, uses 10102-44-0, Nitrogen dioxide, uses 17778-80-2, Atomic oxygen, uses 17778-80-2, Atomic oxygen, uses 19287-45-7, Diborane 25377-83-7, Octene 30637-87-7, Hexanone 32073-03-3, Octyne

(in cleaning of contamination from electron-emissive elements)

L65 ANSWER 14 OF 19 HCA COPYRIGHT 2006 ACS on STN

128:263734 Electrochemical device. Giron, Jean-Christophe (Saint-Gobain Vitrage, Fr.). Eur. Pat. Appl. EP 831360 A1 19980325, 20

pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI. (French). CODEN: EPXXDW.

APPLICATION: EP 1997-402158 19970918. PRIORITY: FR 1996-11392
19960918.

AB The invention concerns an electrochem. device contg. at least 1 substrate, at least 1 elec. conductive layer, at least 1 layer which is electrochem. active and susceptible to reversible ion insertion for such cations at H+, Li+, Na+, Ag+, and an electrolyte. The electrolyte contains at least 1 layer of a material which is essentially an oxide-type mineral for which the ionic conduction can be inhibited or amplified by the incorporation of hydrogenated or nitrated compds., notably nitrides.

IT 7664-38-2, Phosphoric acid, uses

7732-18-5, Water, uses

(electrochem. device applicable to electrochromism and radiation technol. (windows))

RN 7664-38-2 HCA

CN Phosphoric acid (7CI, 8CI, 9CI) (CA INDEX NAME)

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RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
IT
     1333-74-0, Hydrogen, uses 7782-44-7, Oxygen, uses
        (electrochem. device applicable to electrochromism and radiation
        technol. (windows))
RN
     1333-74-0 HCA
    Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
CN
    Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
IC
     ICM G02F001-15
     ICS H01M006-18; G01N027-416; H01M010-36; H01M010-34
CC
     73-11 (Optical, Electron, and Mass Spectroscopy and Other Related
    Properties)
     Section cross-reference(s): 52, 72
     electrochem device electrolyte electrochromism ionic conductor;
ST
    oxide metal electrochromic electrochem device; nitride
     electrochromic electrochem device; proton conductor electrochem
    device electrochromism; lithium conductor electrochem device
    electrochromism; sodium conductor electrochem device
    electrochromism; silver conductor electrochem device
    electrochromism; mineral oxide hydrogenated nitrated electrochem
    device; deposition electrochem device electrochromism;
    plasma CVD electrochem device electrochromism; spray coating
    electrochem device electrochromism; automobile window electrochromic
    electrochem device electrolyte
IT
    Heteropoly acids
        (phosphoric; electrochem. device applicable to
       electrochromism and radiation technol. (windows))
IT
    Vapor deposition process
        (plasma; electrochem. device applicable to
       electrochromism and radiation technol. (windows))
IT
    302-01-2, Hydrazine, uses 1304-76-3, Bismuth sesquioxide, uses
    1310-53-8, Germanium dioxide, uses 1312-43-2, Indium oxide (In2O3)
    1313-96-8, Niobium oxide nb2o5
                                     1313-99-1, Nickel monoxide, uses
    1314-20-1, Thorium dioxide, uses 1314-23-4, Zirconium dioxide,
           1314-35-8, Tungsten trioxide, uses 1314-60-9, Antimony
    uses
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1314-61-0, Tantalum oxide ta2o5 1327-33-9, Antimony 1332-29-2, Tin oxide 7440-57-5, Gold, uses 7631-86-9, Silicon dioxide, uses 7664-38-2, Phosphoric 7722-84-1, Hydrogen peroxide (H2O2), uses **7732-18-5**, Water, uses 9002-98-6, PEI 10024-97-2, Nitrous oxide, uses 11099-11-9, Vanadium oxide 11105-45-6 11118-57-3, Chromium oxide 12055-23-1, Hafnium dioxide 12337-18-7, Tantalum pentoxide hydrate 12586-59-3, Proton 12624-27-0, Rhenium oxide 12645-46-4, Iridium oxide 12712-36-6. Antimony pentoxide hydrate 13463-67-7, Titanium oxide, uses 14701-21-4, Silver(1+), uses 17341-24-1, Lithium 1+, uses 17341-25-2, Sodium(1+), uses 18282-10-5, Tin dioxide 39300-70-4, Lithium nickel oxide 39406-95-6, Cerium titanium oxide 50926-11-9, ITO 56939-16-3, Antimony oxide hydrate 60866-78-6, Tantalum titanium oxide 104812-37-5, Antimony tungsten oxide 202847-02-7, Hydrogen nickel oxide 205312-38-5 205396-60-7

(electrochem. device applicable to electrochromism and radiation technol. (windows))

- L65 ANSWER 15 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 123:154636 A proposed safety assurance method and its application to the fusion experimental reactor. Okazaki, T.; Seki, Y.; Inabe, T.; Aoki, I. (Naka Fusion Research Establishment, Japan Atomic Energy Research Institute, Naka-machi, Naka-gun, Ibaraki-ken, Japan). Fusion Engineering and Design, 30(3), 201-16 (English) 1995
  . CODEN: FEDEEE. ISSN: 0920-3796. Publisher: North-Holland.
- AB Importance categorization and hazard identification methods have been proposed for a fusion exptl. reactor. A parameter, the system index, is introduced in the categorization method. The relative importance of systems with safety functions can be classified by the largeness of the system index and whether or not the system acts as a boundary for radioactive materials. This categorization can be used as the basic principle in detg. structure design assessment, seismic design criteria, etc. For the hazard identification the system time energy matrix is proposed, where the time and spatial distributions of hazard energies are used. This approach is formulated more systematically than an ad-hoc identification of hazard events and it is useful to select design basis events which are employed in the assessment of safety designs.
- IT 1333-74-0, Hydrogen, formation (nonpreparative) (safety assurance method and its application to the fusion exptl. reactors in relation to)
- RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

IT 7732-18-5, Water, reactions

(safety assurance method and its application to the fusion exptl. reactors in relation to)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 7782-44-7, Oxygen, reactions

(safety assurance method and its application to the fusion exptl. reactors in relation to tritium combustion energy)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

CC 71-2 (Nuclear Technology)

fusion reactor safety assurance methodol application; plasma fusion safety assurance methodol application; importance categorization fusion reactor; system index importance categorization fusion reactor; hazard identification fusion reactor

IT Electromagnetic field

Heat

Nuclear energy

Oxidation

Radioactive substances

(safety assurance method and its application to the fusion exptl. reactors in relation to)

IT 1333-74-0, Hydrogen, formation (nonpreparative)

(safety assurance method and its application to the fusion exptl. reactors in relation to)

IT 7732-18-5, Water, reactions

(safety assurance method and its application to the fusion exptl. reactors in relation to)

IT 7782-44-7, Oxygen, reactions

(safety assurance method and its application to the fusion exptl. reactors in relation to tritium combustion energy)

L65 ANSWER 16 OF 19 HCA COPYRIGHT 2006 ACS on STN

113:153202 Silane refunctionalization of radio

frequency glow discharge hydrolyzed

AB

IT

RN

CN

IT

RN

CN

IT

RN

CN

IT

RN

CN

CC

```
ePTFE membrane surfaces. Vargo, T. G.; Hook, D. J.;
     Litwiler, K. S.; Bright, F. V.; Gardella, J. A., Jr. (Univ. Buffalo,
     Buffalo, NY, 14214, USA). Polymeric Materials Science and
     Engineering, 62, 259-63 (English) 1990. CODEN: PMSEDG.
     ISSN: 0743-0515.
     Modified PTFE membranes were functionalized with OH groups
     during treatment with H-H2O or H-MeOH plasma.
     hydrophobicity of the membranes was unaffected by
     plasma treatment. Plasma-treated modified PTFE
     membranes were silvlated with (γ-
     aminopropyl) triethoxysilane.
     7732-18-5, Water, properties
        (plasma contg. hydrogen or oxygen and, PTFE deriv.
        membranes hydrolyzed by, wettability of)
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
H20
     7782-44-7, Oxygen, properties
        (plasma contg. hydrogen or water and, PTFE deriv.
        membranes hydrolyzed by, wettability of)
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
     1333-74-0, Hydrogen, reactions
        (plasma contg., PTFE deriv. membranes
        hydrolyzed by, silylation of)
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-H
     7732-18-5
        (wettability, of plasma-hydrolyzed PTFE deriv.
        membranes, plasma type effect on)
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
     35-8 (Chemistry of Synthetic High Polymers)
```

ST polytetrafluoroethylene deriv membrane plasma treatment; silylation plasma treated polytetrafluoroethylene membrane; wettability PTFE membrane plasma treatment IT Membranes (PTFE derivs., plasma hydrolysis and silylation of, wettability in relation to) IT Solvolysis (of PTFE deriv. membranes, by plasma) IT Silylation (of hydrolyzed PTFE deriv. membranes) IT Wettability (of plasma-hydrolyzed PTFE deriv. membranes, plasma type effect on) IT 919-30-2DP,  $(\gamma$ -Aminopropyl)triethoxysilane, reaction products with hydrolyzed PTFE derivs. 9002-84-0DP, Polytetrafluoroethylene, derivs., hydrolyzed, reaction products with (γaminopropyl)triethoxysilane (membranes, prepn. of, plasma treatment in, membrane wettability in relation to) IT 67-56-1, Methanol, reactions (plasma contg. hydrogen and, PTFE deriv. membranes hydrolyzed by, silylation of) IT 7732-18-5, Water, properties (plasma contg. hydrogen or oxygen and, PTFE deriv. membranes hydrolyzed by, wettability of) 7782-44-7, Oxygen, properties IT (plasma contg. hydrogen or water and, PTFE deriv. membranes hydrolyzed by, wettability of) IT 1333-74-0, Hydrogen, reactions (plasma contg., PTFE deriv. membranes hydrolyzed by, silylation of) IT 7732-18-5 (wettability, of plasma-hydrolyzed PTFE deriv. membranes, plasma type effect on) ANSWER 17 OF 19 HCA COPYRIGHT 2006 ACS on STN 110:146878 Development and application of a microwave-induced plasma emission spectrometric detector for gas chromatography in China. Yu, Weile (Lanzhou Inst. Chem. Phys., Acad. Sin., Lanzhou, Peop. Rep. China). Journal of Analytical Atomic Spectrometry, 3(6), 893-900 (English) 1988.

JASPE2. ISSN: 0267-9477.

AB Two Chinese-built gas chromatog./microwave-induced plasma emission spectrometry (GC-MIP-ES) systems (Model SG-1) and one imported system (Model MPD-850) were used at the Lanzhou Institute of Chem. Physics, Academia Sinica (LICP), the Shanghai Institute of Org. Chem., Academia Sinica (SIOC), and the Research Institute of

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H<sub>2</sub>O

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Alcohols, analysis

the Shanghai Petroleum Refinery of the Gao Qiao Petrochem. Corporation (SPR). A diverse range of research applications (such as basic chem. reactions, industrial chem. analyses, and methodol. studies) were investigated at these three institutes. Various selected examples are cited to demonstrate the characteristics of the GC-MIP-ES system. **7732-18-5**, Water, analysis (detn. of, in Freon-12 by gas chromatog. with microwave-induced plasma emission spectrometric detector) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 1333-74-0, Hydrogen, analysis 7553-56-2, Iodine, analysis 7782-44-7, Oxygen, analysis (detn. of, in org. compds. by gas chromatog. with microwave-induced plasma emission spectrometric detector) 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7553-56-2 HCA Iodine (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 080-2 (Organic Analytical Chemistry) Section cross-reference(s): 79 gas chromatog plasma emission spectrometric detector; microwave induced plasma emission spectrometric detector; China plasma spectrometer gas chromatog detector Chemical formula (detn. of, by gas chromatog. with microwave-induced plasma emission spectrometric detector)

(detn. of, in gasoline by gas chromatog. with plasma

emission spectrometric detector)

- IT Chromatographs, gas
  - (detectors, spectrometric, with microwave-induced plasma
- IT 60320-18-5, Tetrachloropropene
  - (anal. of raw, by gas chromatog. with plasma emission spectrometric detector)
- TT 75-43-4, Dichlorofluoromethane 75-45-6, Chlorodifluoromethane 75-69-4, Trichlorofluoromethane 75-72-9, Chlorotrifluoromethane 124-38-9, Carbon dioxide, analysis 7732-18-5, Water, analysis
  - (detn. of, in Freon-12 by gas chromatog. with microwave-induced plasma emission spectrometric detector)
- IT 593-74-8, Dimethyl mercury 29138-86-1, Ethylmethyl mercury (detn. of, in di-Et mercury by gas chromatog. with microwave-induced plasma emission spectrometric detector)
- IT 64-17-5, Ethanol, analysis 71-36-3, 1-Butanol, analysis 75-65-0, Tert-Butanol, analysis 78-92-2, 2-Butanol (detn. of, in gasoline by gas chromatog. with plasma emission spectrometric detector)
- IT **1333-74-0**, Hydrogen, analysis 7439-97-6, Mercury, analysis 7440-44-0, Carbon, analysis **7553-56-2**, Iodine, analysis
  - 7704-34-9, Sulfur, analysis 7723-14-0, Phosphorus, analysis
  - 7726-95-6, Bromine, analysis 7727-37-9, Nitrogen, analysis
  - 7782-39-0, Deuterium, analysis 7782-41-4, Fluorine, analysis
  - 7782-44-7, Oxygen, analysis 7782-50-5, Chlorine, analysis (detn. of, in org. compds. by gas chromatog. with microwave-induced plasma emission spectrometric detector)
- TT 78-87-5, 1,2-Dichloropropane 96-18-4, 1,2,3-Trichloropropane 96-19-5 142-28-9, 1,3-Dichloropropane 1070-78-6 3175-23-3 10436-39-2, 1,1,2,3-Tetrachloro-1-propene 16500-91-7 (detn. of, in raw tetrachloropropene by gas chromatog. with plasma emission spectrometric detector)
- IT 119797-78-3
  - (deuterium detn. in, and theor. empirical formula calcn. for, by gas chromatog. with **plasma** emission spectrometric detection)
- IT 67-56-1, Methanol, analysis 67-64-1, Acetone, analysis 71-43-2, Benzene, analysis 95-47-6, analysis 95-63-6, 1,2,4-Trimethylbenzene 141-78-6, Ethyl acetate, analysis 142-82-5, Heptane, analysis 98-06-6, Tert-Butylbenzene 100-41-4, 108-38-3, m-Xylene, analysis Ethylbenzene, analysis 108-67-8, 108-87-2, Methyl cyclohexane 108-88-3, analysis 111-65-9, Octane, analysis 110-82-7, Cyclohexane, analysis 111-84-2, Nonane 124-18-5, Decane 1120-21-4, Undecane

(identification of, in gasoline by gas chromatog. with

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plasma emission spectrometric detector)
IT
                  82949-11-9
                               118992-99-7
     82949-08-4
        (identification of, in reaction mixts. by gas chromatog, with
        plasma emission spectrometric detector)
IT
     75-71-8, Freon-12 627-44-1, Diethyl mercury
        (impurity detn. in, by gas chromatog. with microwave-induced
        plasma emission spectrometric detector)
    ANSWER 18 OF 19 HCA COPYRIGHT 2006 ACS on STN
97:207384 Techniques for quantitation and identification of organic
     micropollutants by high resolution gas chromatography and element
     specific emission spectroscopy. Stieglitz, L.; Zwick, G. (Inst.
     Heisse Chem., Kernforschungszent., Karlsruhe, Fed. Rep. Ger.).
     Comm. Eur. Communities, [Rep.] EUR, EUR 7623, Anal. Org.
     Micropollut. Water, 105-12 (German) 1982. CODEN: CECED9.
AΒ
     A combination of a gas chromatog. and a microwave-plasma
     emission detector was modified for use with high-resoln. capillary
     columns for detn. and identification of org. micropollutants.
     Eleven elements (C, H, D, N, P, O, S, F, Cl, Br, and I) can be
     monitored simultaneously in the gas chromatog. effluents.
     relative molar response of the elements in the compds. varies only
     by .apprx. ±10%. The detection limits were in the range of ng/s.
     From the elemental ratios measured, the identification of the
     various compds. was possible. The technique was illustrated by
     identification of organohalogen compds. in water samples.
IT
     1333-74-0, analysis 7553-56-2, analysis
     7782-44-7, analysis
        (detn. of, in org. compds. by capillary gas chromatog. with
        plasma emission spectrometric detection)
RN
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7553-56-2 HCA
CN
     Iodine (8CI, 9CI) (CA INDEX NAME)
I-I
     7782-44-7 HCA
RN
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
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7732-18-5, analysis

IT

(organohalogen pollutant identification in, by capillary gas chromatog. with plasma emission spectrometric detection) 7732-18-5 HCA RN · CN Water (8CI, 9CI) (CA INDEX NAME) H<sub>2</sub>O CC 80-6 (Organic Analytical Chemistry) Section cross-reference(s): 61 STorg pollutant detn chromatog spectroscopy; gas chromatog org pollutant detn; emission spectroscopy chromatog pollutant detn; plasma spectroscopy chromatog pollutant detn; water analysis organohalogen chromatog spectroscopy ITChromatography, gas (capillary, with plasma emission spectrometric detection, for detn. or identification of org. pollutants) IT Spectrochemical analysis (emission, with plasma source, gas chromatog. combined with, for detn. and identification of org. pollutants) Halogen compounds IT (org., identification of, in water by capillary gas chromatog. with plasma emission spectrometric detection) IT 87-68-3 100-68-5 108-86-1, analysis 110-02-1 118-74-1 462-06-6 344-03-6 352-32-9 443-26-5 541-73-1 544-76-3 616-44-4 705-29-3 827-08-7 1002-69-3 1146-65-2 3114-55-4 4165-57-5 4549-32-0 41051-88-1 (detn. of, by capillary gas chromatog. with plasma emission spectrometric detection) IT 1333-74-0, analysis 7440-44-0, analysis 7553-56-2 , analysis 7704-34-9, analysis 7723-14-0, analysis 7726-95-6, analysis 7727-37-9, analysis 7782-39-0, analysis 7782-41-4, analysis 7782-44-7, analysis 7782-50-5, analysis (detn. of, in org. compds. by capillary gas chromatog. with plasma emission spectrometric detection) IT **7732-18-5**, analysis (organohalogen pollutant identification in, by capillary gas chromatog. with plasma emission spectrometric detection) ANSWER 19 OF 19 HCA COPYRIGHT 2006 ACS on STN 83:85995 Thermodynamic properties of plasma forming gases. Suris, A. L.; Aslanyan, L. S.; Shorin, S. N. (Mosk. Inst. Khim. Mashinostr., Moscow, USSR). Khimiya Vysokikh Energii, 8(5), 392-4 (Russian) 1974. CODEN: KHVKAO. ISSN: 0023-1193.

Thermodn. calcns. of the equil. state of plasma-forming

AB

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RN

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H-H

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I-I

RN

CN

H<sub>2</sub>O

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CC

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IT

IT

IT

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gases H, O, N, S, F, Cl, Br, I, H2S, H2O, HCl, HF, HBr, HI, CH4,
     CF4, SO2, CO2, and NH3 at 293-6000°K and 0.1-30 bar show that
     CH4 and CF4 require the highest amt. of energy for plasma
                 The highest sound-propagation velocity was exhibited by
     formation.
     H and those gases which on dissocn. form H.
     1333-74-0, properties 7553-56-2, properties
     7732-18-5, vapor 7782-44-7, properties
        (plasma, thermal dissocn. of and sound velocity in)
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
     7553-56-2 HCA
     Iodine (8CI, 9CI) (CA INDEX NAME)
     7732-18-5 HCA
    Water (8CI, 9CI) (CA INDEX NAME)
     7782-44-7 HCA
    Oxygen (8CI, 9CI)
                      (CA INDEX NAME)
0===0
     69-2 (Thermodynamics, Thermochemistry, and Thermal Properties)
     Section cross-reference(s): 68, 65
    plasma thermodn calcn gas; energy loss gas plasma
    Thermodynamics
        (of thermal plasma formation, in gases)
    Dissociation
        (thermal, of plasma-forming gases)
    Plasma
        (thermodn. of gases forming)
    Sound and Ultrasound
        (velocity of, in plasma-forming gases)
    74-82-8, properties 75-73-0 124-38-9, properties
    1333-74-0, properties 7446-09-5, properties
    7553-56-2, properties
                            7664-41-7, properties 7704-34-9,
                 7726-95-6, properties
                                        7727-37-9, properties
    properties
    7732-18-5, vapor 7782-41-4, properties 7782-44-7
```

, properties 7782-50-5, properties 7783-06-4, properties (plasma, thermal dissocn. of and sound velocity in)

IT 7647-01-0, properties 7664-39-3, properties 10034-85-2 10035-10-6, properties (plasma, thermal dissocn. of and sound velocity in gaseous)

## => D L66 1-32 CBIB ABS HITSTR HITIND

L66 ANSWER 1 OF 32 HCA COPYRIGHT 2006 ACS on STN
140:170861 High performance tokamak experiments with a ferritic steel
 wall on JFT-2M. Tsuzuki, K.; Kimura, H.; Kawashima, H.; Sato, M.;
 Kamiya, K.; Shinohara, K.; Ogawa, H.; Hoshino, K.; Bakhtiari, M.;
 Kasai, S.; Uehara, K.; Sasao, H.; Kusama, Y.; Isei, N.; Miura, Y.;
 Ogawa, T.; Yamamoto, M.; Shibata, T.; Kikuchi, K.; Miyachi, K.; Ito,
 T.; Ajikawa, H.; Nomura, S.; Tsutsui, H.; Shimada, R.; Ido, T.;
 Hamada, Y.; Fukumoto, N.; Nagata, M.; Uyama, T.; Niimi, H.; Yatsu,
 S.; Kayukawa, N.; Hino, T.; Hirohata, Y.; Nagashima, Y.; Ejiri, A.;
 Amemiya, A.; Sadamoto, Y.; Tsushima, A. (Naka Fusion Research
 Establishment, Japan Atomic Energy Research Institute, Tokai-mura,
 319-1195, Japan). Nuclear Fusion, 43(10), 1288-1293 (English)
 2003. CODEN: NUFUAU. ISSN: 0029-5515. Publisher:
 Institute of Physics Publishing.

AB Compatibility between the plasma and low activation ferritic steel, which is a candidate material for fusion demonstration reactors, has been investigated step by step in the JFT-2M tokamak. The authors have entered the third stage of the Advanced Material Tokamak Expt. (AMTEX), where the inside of the vacuum vessel wall is completely covered with ferritic steel plates ferritic inside wall (FIW). The effects of a FIW on the plasma prodn., impurity release, the operation region, and H-mode characteristics have been investigated. No neg. effect has been obsd. up to now. A high normalized beta plasma of βN .apprx. 3, having both an internal transport barrier and a steady H-mode edge was obtained. A remarkable redn. in ripple trapped loss from 0.26 MW m-2 (without ferritic steel) to less than 0.01 MW m-2 was demonstrated by the optimization of the thickness profile of FIW. A code to calc. fast ion losses, taking into account the full 3-dimensional magnetic structure was developed, and values obtained using the code showed good agreement with exptl. Thus, encouraging results are obtained for the use of this material in the DEMO-reactor.

IT 7782-44-7, Oxygen, processes

RN

(impurity; tokamak expts. with a ferritic steel wall on JFT-2M) 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0==0

IT 7732-18-5, Water, processes

(tokamak expts. with a ferritic steel wall on JFT-2M and assocd. cleaning issues)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0, Hydrogen, processes

(tokamak expts. with a ferritic steel wall on JFT-2M and assocd. cleaning issues)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

CC 71-2 (Nuclear Technology)
 Section cross-reference(s): 55

ST tokamak fusion plasma ferritic steel wall expt JFT2M; impurity ferromagnetic effect ferritic steel wall tokamak

IT 7782-44-7, Oxygen, processes

(impurity; tokamak expts. with a ferritic steel wall on JFT-2M)

IT 124-38-9, Carbon dioxide, processes 630-08-0, Carbon monoxide, processes 7732-18-5, Water, processes

(tokamak expts. with a ferritic steel wall on JFT-2M and assocd. cleaning issues)

IT 1333-74-0, Hydrogen, processes

(tokamak expts. with a ferritic steel wall on JFT-2M and assocd. cleaning issues)

IT 7440-59-7, Helium, uses

(tokamak expts. with a ferritic steel wall on JFT-2M and assord. glow discharge cleaning)

L66 ANSWER 2 OF 32 HCA COPYRIGHT 2006 ACS on STN

139:263096 Production of hydrogen from a hydrocarbon fuel. Bowe,
Michael Joseph; Hall, Stephen Ivor; Martin, Anthony Robert (Accentus
PLC, UK). PCT Int. Appl. WO 2003078308 A2 20030925, 23

pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG,

BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES,

FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,

KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR,

TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW; RW: AT, BE, BF, BJ,

CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2003-GB978 20030307. PRIORITY: GB 2002-5837 20020313; GB 2002-15414 20020704.

Hydrogen is produced from a hydrocarbon fuel, such as diesel, by AB subjecting a mixt. of the fuel with oxygen gas to plasma treatment in a dielec. barrier plasma reactor to generate oxygenated mols.; mixing the resulting oxygenated mols. with steam and subjecting them to steam reforming in a compact catalytic reactor at 550-850°, and then to a water gas shift reaction (possibly with addnl. steam) at 500-700°. The source of heat for the endothermic reactions is provided by catalytic combustion in the flow channels. The resulting gases may then be mixed with a small quantity of oxygen gas, and subjected to selective oxidn. to convert any carbon monoxide to carbon dioxide. The oxygen is produced by electrolysis of water. The hydrogen is sepd. from the other products using a hydrogen-permeable membrane. hydrogen may be subsequently used in a fuel cell to generate electricity. The produced CO2 is injected into porous rock formation below the sea bed. Such a process may be used at an offshore facility.

IT 1333-74-0P, Hydrogen, uses

(prodn. of hydrogen from hydrocarbon fuel)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

TT 7732-18-5, Water, reactions 7782-44-7, Oxygen,
reactions

(prodn. of hydrogen from hydrocarbon fuel)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IC ICM C01B003-02

CC 51-11 (Fossil Fuels, Derivatives, and Related Products) Section cross-reference(s): 52, 67

ST hydrogen manuf purifn fuel cell power generation; plasma oxidn steam reforming water gas shift reaction hydrogen IT Reactors (plasma; prodn. of hydrogen from hydrocarbon fuel) IT Diesel fuel Electrolysis Fuel cells Membranes, nonbiological Power generation Steam reforming catalysts Water gas shift reaction Water gas shift reaction catalysts (prodn. of hydrogen from hydrocarbon fuel) IT Fuel gas manufacturing (steam reforming; prodn. of hydrogen from hydrocarbon fuel) 1333-74-0P, Hydrogen, uses IT (prodn. of hydrogen from hydrocarbon fuel) 7732-18-5, Water, reactions 7782-44-7, Oxygen, IT reactions (prodn. of hydrogen from hydrocarbon fuel) ANSWER 3 OF 32 HCA COPYRIGHT 2006 ACS on STN L66 139:189022 Post metal etch polymer removal: A new CF4-based dry plasma process sequence. Pozzoli, Maria Paola; Petroni, Simona (Novellus Systems, Inc., San Jose, CA, 95134, USA). Diffusion and Defect Data -- Solid State Data, Pt. B: Solid State Phenomena, 92 (Ultra Clean Processing of Silicon Surfaces V), 255-258 (English) 2003. CODEN: DDBPE8. ISSN: 1012-0394. Publisher: Scitec Publications. AB A novel CF4 based method for the dry plasma removal of post-metal etch polymer residues has been developed on a dual plasma source, dry cleaning system. This process eliminates the via damage problems obsd. when the traditional wet solvent clean process is used on specific embedded flash devices. In order to effectively clean these products, it was found that the CF4/O2/N2/H2 MW and RF dry plasma process needs to be followed by an intermediate H2O rinse (either deionized water or in-situ H2O vapor within the plasma reactor), then by a second CF4/N2H2 RF dry plasma step, and by a final H2O rinse. The intermediate H2O step appears

to be crucial in making the fluorinated polymer residues more sol.,

process provides a reliable and cost-effective alternative method to solvent-based processing, with greater cleaning efficiency, less

The above

before their exposure to the second F-based dry plasma step and their final elimination through H2O rinse. The

damage to device structures and less environmental impacts.

1333-74-0, Hydrogen, processes 7782-44-7, Oxygen, IT processes (etchant; post metal etch polymer removal using CF4-based dry plasma process sequence) 1333-74-0 HCA RNCN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CNo = oIT 7732-18-5, Water, processes (rinsing; post metal etch polymer removal using CF4-based dry plasma process sequence) RN 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H20 CC 76-3 (Electric Phenomena) Section cross-reference(s): 38 IT (plasma; post metal etch polymer removal using CF4-based dry plasma process sequence) IT Semiconductor device fabrication (post metal etch polymer removal using CF4-based dry plasma process sequence) IT Fluoropolymers, processes (post metal etch polymer removal using CF4-based dry plasma process sequence) IT Washing (rinsing; post metal etch polymer removal using CF4-based dry plasma process sequence) IT 75-73-0, Tetrafluoromethane 1333-74-0, Hydrogen, processes 7727-37-9, Nitrogen, processes 7782-44-7, Oxygen, processes (etchant; post metal etch polymer removal using CF4-based dry plasma process sequence) 7440-21-3, Silicon, processes 7440-50-8, Copper, processes IT (post metal etch polymer removal using CF4-based dry plasma process sequence) IT 7732-18-5, Water, processes

(rinsing; post metal etch polymer removal using CF4-based dry plasma process sequence)

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ANSWER 4 OF 32 HCA COPYRIGHT 2006 ACS on STN
136:223943 Procedure and device for the frequency-modulated excitation
     of hydrogen and/or a water vapour air mixture
     plasma with high frequency oscillations. Zembold, Otmar
     (Nanoplasmon Gmbh, Germany). Ger. Offen. DE 10041065 A1
     20020307, 6 pp. (German). CODEN: GWXXBX. APPLICATION: DE
     2000-10041065 20000822.
     This high frequency oscillator produces ionized hydrogen or
AΒ
     water vapor. The device consists of a specialized
     chamber in a suitable contq. vessel, made from dielec. material,
     with the gas that will be ionized. Self-induction and a condenser
     creates a resonant circuit in the vessel and they are arranged in a
     manner so that the resonance frequency F0 is in the high frequency
     range and that the open circuit overvoltage coeff., Q0, is defined
     by Q0 = 2\pi F0L/R = 1/(2\pi F0CR). In this equation L =
     self-induction parameter, C = condenser capacity, and R = resonant
     circuit resistance. Since Q0>1, the resonance condition produces a
     trigger- or excitation voltage in the condenser, which lead to
     ionization of the gas by the electromagnetic radiation which has a
     frequency of \approx F0. Construction details are presented.
IT
     1333-74-0, Hydrogen, processes 7732-18-5, Water,
     processes 7782-44-7, Oxygen, processes
        (frequency-modulated ionization of hydrogen and/or water
        vapor-air plasma with high frequency
        oscillations)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-- H
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H<sub>2</sub>O
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
```

IC ICM H05H001-46 ICS G01N021-71 CC 73-10 (Optical, Electron, and Mass Spectroscopy and Other Related Properties)

Section cross-reference(s): 65, 76

- ST frequency modulated excitation hydrogen water plasma microwave oscillator resonator
- IT Electric insulators
  Energy level excitation
  Masers

Microwave oscillators
Overvoltage

Water vapor

(frequency-modulated ionization of hydrogen and/or water vapor-air plasma with high frequency oscillations)

IT Resonators

(microwave; frequency-modulated ionization of hydrogen
and/or water vapor-air plasma with
high frequency oscillations)

- 1333-74-0, Hydrogen, processes 7732-18-5, Water,
   processes 7782-44-7, Oxygen, processes
   (frequency-modulated ionization of hydrogen and/or water
   vapor-air plasma with high frequency
   oscillations)
- L66 ANSWER 5 OF 32 HCA COPYRIGHT 2006 ACS on STN

  136:190585 Three-dimensional analysis of combustible mixture generation in an ITER-FEAT first-wall coolant leak scenario. Baumann, W.;
  Breitung, W.; Kaup, B.; Necker, G.; Royl, P.; Travis, J. R.
  (Programm Kernfusion, Inst. Kem- Energietechnik, Forschungszentrum Karlsruhe GmbH, Karlsruhe, Germany). Wissenschaftliche Berichte Forschungszentrum Karlsruhe, FZKA 6663, i-v, 1-37 (English)

  2001. CODEN: WBFKF5. ISSN: 0947-8620.
- Three-dimensional distribution calcns. are performed for the AB ITER-FEAT vacuum vessel, the connected pressure suppression pool and drain tank. An ex-vessel/first-wall coolant leak without plasma shutdown is simulated. The steam, H, and air sources for this sequence are taken from best-est. MELCOR The time- and space-dependent gas distribution in the system is calcd. using the verified 3-dimensional Computational Fluid Dynamics code GASFLOW. A new extended version of GASFLOW was developed to model the ITER-FEAT specific phenomena in adequate detail. During the accident sequence, H initially appears only in the vacuum vessel due to the steam/Be reaction. failure of the burst membranes, steam and H flow from the vacuum vessel through the connecting lines to the suppression pool and the drain tank. Because of the ongoing steam condensation occurring in the suppression pool, the pressure there remains permanently at a lower level compared to the

other components, resulting in a continuous flow of steam and noncondensable gases into this vol. Since no steam condensation is modeled in the drain tank, almost all H2 and N2 accumulate in the suppression pool cover gas vol. After 10,500 s of steam flow, also air starts entering the vacuum vessel, with the basic mechanisms remaining the same. Consequently, an accumulation of N2 and O2 takes place in the suppression pool cover gas. Combustible and explosive H2-O2-N2 mixts. exist after 13,600 s, and at 21,000 s a stoichiometric H2/O2 ratio has formed, involving 14 kg of H. Contrary to the situation in the suppression pool with its significant H risk, only inert, steam dominated mixts. without hazard potential develop in the vacuum vessel and the drain tank. Various passive mitigation measures could be considered to reduce or completely remove the H risk in the suppression pool.

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

CC 71-2 (Nuclear Technology)

IT Steam

(continuous flow of **steam** and noncondensable gases in suppression pool happens)

1333-74-0, Hydrogen, formation (nonpreparative) 7727-37-9, Nitrogen, formation (nonpreparative) 7782-44-7, Oxygen, formation (nonpreparative)

(combustible and explosive H2-O2-N2 mixts. exist)

IT 7440-41-7, Beryllium, reactions

(hydrogen initially appears during accident in vacuum vessel due to **steam**/beryllium reaction.)

L66 ANSWER 6 OF 32 HCA COPYRIGHT 2006 ACS on STN

136:160223 Magnetic recording medium and method for producing and inspecting the deposition characteristics in its fabrication. Kato Junya; Suzuki, Mikio (Showa Denko K.K., Japan). PCT Int. Appl. WO 2002011130 Al 20020207, 93 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR,

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CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-JP6580 20010731. PRIORITY: JP 2000-231114 20000731; US 2000-246595P 20001108.
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AB The present invention provides a method for inspecting deposition characteristics of a deposition on the surface of a protective film predominantly contg. C of a magnetic recording medium, which medium includes a disk and the protective film formed on the disk, the disk including a nonmagnetic substrate, a nonmagnetic undercoat layer, and a magnetic layer, the layers being formed on the substrate, wherein the method includes comparing a predetd. threshold with the extn. amt. of an inspection gas component and/or a compd. component formed so as to contain the inspection gas component, the gas component and/or the compd. component being extd. with an inspection solvent after the magnetic recording medium is allowed to stand in an atm. of the inspection gas component; a process for producing a magnetic recording medium in which the extn. amt. is or greater than the threshold, the extn. amt. being obtained through the inspection method; and a magnetic recording medium produced through the prodn. process.

IT 1333-74-0, Hydrogen, uses

(amorphous carbon dopant; magnetic recording medium and method for producing and inspecting deposition characteristics in fabrication)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

## IT 7732-18-5, Water, processes

(inspection solvent; magnetic recording medium and method for producing and inspecting deposition characteristics in fabrication)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

## IT 7782-44-7, Oxygen, processes

(magnetic recording medium and method for producing and inspecting deposition characteristics in fabrication)

RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME)

o = 0

- IC ICM G11B005-72 ICS G11B005-84
- CC 77-8 (Magnetic Phenomena)

- IT 64-17-5, Ethanol, processes 67-56-1, Methanol, processes 67-63-0, Isopropanol, processes **7732-18-5**, Water, processes
  - (inspection solvent; magnetic recording medium and method for producing and inspecting deposition characteristics in fabrication)
- ANSWER 7 OF 32 HCA COPYRIGHT 2006 ACS on STN Method for production of enhanced traceable immunizing 135:362523 drinking water and other liquid and gas products, devices for production and use thereof, and use of the enhanced products for immunizing living beings. Tribelsky, Zamir; Ende, Michael (Atlantium Ltd., Israel). PCT Int. Appl. WO 2001083385 A2 20011108, 139 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-IL383 20010427. PRIORITY: IL 2000-135843 20000428.

AB A method for the prodn. of enhanced traceable optp-physiol. polished liqs., and gases or solids or combination for immunizing living beings, devices using the method, use, and preferred mode for utilization are disclosed. A multi processing platform is proposed according to the invention harnessing time domain optronics of light and sound, wherein the transient sound produced by light is measured, referenced or calibrated against the light produced by sound for the formation adequate energy levels or densities or fluence rates for the purpose of dissocn. of noxious or innocuous species or combination constituents components while keeping their geometrical integrity above their predetd. resonance levels, thus intact for later traceable recognition and triggering of pos. decisive action by immune systems.

IT 7782-44-7, Oxygen, biological studies

(dissolved; prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

IT 7732-18-5, Water, reactions

(oxidn., supercrit.; prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0, Hydrogen, biological studies

(prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

IC ICM C02F009-00

ICS B01D053-00

CC 63-3 (Pharmaceuticals)

Section cross-reference(s): 15, 16, 17, 18, 47, 61

IT Cell membrane

(components; prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)

- IT Plasma
  - (non-thermal; prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)
- IT Lymphocyte
  - (plasma cell; prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)

- IT 1333-74-0, Hydrogen, biological studies (prodn. of biol. enhanced traceable drinking water and other liq. and gas products for immunizing living beings)
- L66 ANSWER 8 OF 32 HCA COPYRIGHT 2006 ACS on STN

  135:275384 Mixed reactant fuel cells with flow through porous electrodes. Priestnall, Michael Alexander; Evans, Michael Joseph; Shaffer, Milo Sebastian Peter (Scientific Generics Limited, UK).

  PCT Int. Appl. WO 2001073881 Al 20011004, 68 pp.

  DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English).

  CODEN: PIXXD2. APPLICATION: WO 2001-GB1339 20010326. PRIORITY: GB
- AB A fuel cell or battery for providing useful elec. power by electrochem. means, comprises: at least one cell; at least one anode and at least one cathode within the cell, and ion-conducting electrolyte means for transporting ions between the electrodes; and is characterized in that: the electrodes are porous and in that means are provided for causing hydrodynamic flow of a mixt. of at least fuel and oxidant through the body of the electrodes.

GB 2000-25030 20001012; GB 2000-26935 20001103; GB 2000-27587

- IT **7732-18-5**, water, uses
  - (mixed reactant fuel cells with flow through porous electrodes)

2000-7306 20000324; GB 2000-19622 20000809; GB 2000-19623 20000809;

RN 7732-18-5 HCA

20001110.

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O IT **7782-44-7**, Oxygen, reactions (mixed reactant fuel cells with flow through porous electrodes) RN7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0==0 1333-74-0, Hydrogen, uses IT (mixed reactant 'fuel cells with flow through porous electrodes) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CNH-HIC ICM H01M008-10 ICS H01M008-08; H01M008-22 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC ΙT Air Battery electrodes Fuel cell electrodes Fuel cells Plasma Secondary batteries Turbulence (mixed reactant fuel cells with flow through porous electrodes) Membranes, nonbiological IT (polymer; mixed reactant fuel cells with flow through porous electrodes) IT 1310-58-3, potassium hydroxide, uses 7732-18-5, water, uses 7783-96-2, Silver iodide 55575-02-5, Cerium gadolinium 64417-98-7, Yttrium zirconium oxide 65453-23-8, Cerium 65983-06-4, Iridium zirconium oxide zirconium oxide (mixed reactant fuel cells with flow through porous electrodes) 7722-84-1, Hydrogen peroxide, reactions 7782-44-7, Oxygen, IT (mixed reactant fuel cells with flow through porous electrodes) 67-56-1, Methanol, uses 1333-74-0, Hydrogen, uses IT 7440-44-0, Carbon, uses (mixed reactant fuel cells with flow through porous electrodes)

HCA COPYRIGHT 2006 ACS on STN

135:36421 Decomposition of CH3SH in an RF Plasma

ANSWER 9 OF 32

L66

Reactor: Reaction Products and Mechanisms. Tsai, Cheng-Hsien; Lee, Wen-Jhy; Chen, Chuh-Yung; Liao, Wei-Tung (Departments of Environmental Engineering and Chemical Engineering, National Cheng Kung University, Tainan, 70101, Taiwan). Industrial & Engineering Chemistry Research, 40(11), 2384-2395 (English) 2001. CODEN: IECRED. ISSN: 0888-5885. Publisher: American Chemical Society.

AB Application of the RF (radio frequency ) cold plasma method to the decompn. of methanethiol (Me mercaptan, CH3SH) at different O2/CH3SH ratios (0-4.5), with various input powers (20-90 W), and at const. operating pressure (30 Torr) was investigated. The species detected in the CH3SH/O2/Ar RF plasma were SO2, CS2, OCS, CO, CO2, CH4, C2H4 C2H2, H2, H2O, HCOH, and CH3OH. However, CS2, CH4, C2H4, C2H2, H2, H2S, CH3SCH3 (DMS), and CH3S2CH3 (DMDS) were detected in the CH3SH/Ar RF plasma. In the CH3SH/Ar plasma, over 83.7% of the total sulfur input was converted into CS2 at 60 W; this is due to the lack of competition between O and S and the thermodn. stability of CS2. In the oxygen-rich conditions of the CH3SH/O2/Ar plasma, the most predominant sulfur-contg. compd. was SO2. As the feed O2/CH3SH ratio was increased, MSO2 was increased, while MCS2 was decreased simultaneously. MOCS was reduced by increasing either the O2/CH3SH ratio or the applied power. From the decay of CS2 and the generation of CO at a lower O2/CH3SH ratio of 0.6, CS, CS2, and CO were suggested as the primary species to react with O, OH, O2, S, or This study provides useful insight into S2 and then to form OCS. the reaction mechanisms involved in the decompn. of CH3SH and, mainly, the formation of CS2, CH4, C2H2, C2H4, SO2, CO, CO2, and OCS. in CH3SH/Ar and CH3SH/O2/Ar plasmas.

IT 7732-18-5, Water, processes

(reaction product of decompn. of CH3SH in CH3SH/O2/Ar mixt. in radio frequency plasma reactor)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0, Hydrogen, processes

(reaction product of decompn. of CH3SH in both CH3SH/O2/Ar and CH3SH/Ar mixts. in radio frequency

plasma reactor)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

IT 7782-44-7, Oxygen, reactions (reaction products and mechanisms of decompn. of CH3SH in radio frequency plasma reactor) RN

7782-44-7 HCA

Oxygen (8CI, 9CI) (CA INDEX NAME) CN

0 = 0

CC 59-2 (Air Pollution and Industrial Hygiene)

ST mechanism decompn methanethiol radio frequency plasma; odor control mechanism decompn methanethiol radio frequency plasma

Air pollution IT

> (control, odor; reaction products and mechanisms of decompn. of CH3SH in radio frequency plasma reactor and implications for odor control)

IT Cold plasma

Decomposition kinetics

Reaction mechanism

(reaction products and mechanisms of decompn. of CH3SH in radio frequency plasma reactor)

IT 7440-37-1, Argon, miscellaneous (inert gas used to study reaction products and mechanisms of decompn. of CH3SH in radio frequency plasma reactor)

IT 75-18-3, Dimethylsulfide 624-92-0, Dimethyldisulfide 7783-06-4, Hydrogen sulfide, processes (reaction product of decompn. of CH3SH in CH3SH/Ar mixt. in

radio frequency plasma reactor) 50-00-0, Formaldehyde, processes 67-56-1, Methanol, processes IT 124-38-9, Carbon dioxide, processes 463-58-1, Carbon oxide sulfide 630-08-0, Carbon monoxide, processes 7446-09-5, Sulfur dioxide, processes 7732-18-5, Water, processes

> (reaction product of decompn. of CH3SH in CH3SH/O2/Ar mixt. in radio frequency plasma reactor)

74-82-8, Methane, processes IT 74-85-1, Ethene, processes 74-86-2, Ethyne, processes 75-15-0, Carbon disulfide, processes 1333-74-0, Hydrogen, processes

> (reaction product of decompn. of CH3SH in both CH3SH/O2/Ar and CH3SH/Ar mixts. in radio frequency plasma reactor)

74-93-1, Methanethiol, processes IT (reaction products and mechanisms of decompn. of CH3SH in an **RF plasma** reactor)

ΙŢ 7782-44-7, Oxygen, reactions (reaction products and mechanisms of decompn. of CH3SH in

## radio frequency plasma reactor)

ST

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L66 ANSWER 10 OF 32 HCA COPYRIGHT 2006 ACS on STN
135:11229 Removal of particles by ICRF cleaning in HT-7 superconducting
     tokamak. Hu, Jian-sheng; Li, Jian-gang; Zhang, Shou-yin; Gu,
     Xue-mao; Zhang, Xiao-dong; Zhao, Yan-pin; Gong, Xian-zu; Kuang,
     Guang-li; Li, Cheng-fu; Luo, Jia-rong; Wang, Xiao-ming; Gao, Xiang;
     Wan, Bao-nian; Xie, Ji-kang; Wan, Yuan-xi (Institute of Plasma
     Physics, The Chinese Academy of Sciences, Hefei, 230031, Peop. Rep.
     China). Plasma Science & Technology (Hefei, China), 3(1), 621-627
     (English) 2001. CODEN: PSTHC3.
                                      ISSN: 1009-0630.
     Publisher: Chinese Academy of Sciences, Institute of Plasma Physics.
AB
     The ICRF (Ion Cyclotron Range Frequency) cleaning technique has been
     used as a routine wall cleaning method in the HT-7 superconducting
     tokamak. In a wide range of toroidal fields, the removal rate of
     residual gas by ICRF cleaning was about 20-fold higher than that of
     glow discharge cleaning (GDC). At different gas
     pressures and RF power levels, the ICRF cleaning is
     studied carefully. A good impurity cleaning effect and a very high
     hydrogen removal rate were obtained. The removal rate of hydrogen
     by 5 kW ICRF cleaning achieved was 1.6 + 10-5 torr L/s.
     Relations among pressure, outgassing rate, at. layers absorbed on
     surface and the cleaning mode are discussed.
     1333-74-0, Hydrogen, processes 7732-18-5, Water,
IT
     processes 7782-44-7, Oxygen, processes
        (particle removal by ICRF cleaning in the HT-7 superconducting
        tokamak)
     1333-74-0 HCA
RN
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
н-н
RN ·
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H<sub>2</sub>O
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
0 = 0
CC
     71-2 (Nuclear Technology)
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tokamak ICRF cleaning particle removal HT7; fusion plasma

ICRF cleaning particle removal HT7

L66 ANSWER 11 OF 32 HCA COPYRIGHT 2006 ACS on STN
134:358568 Conditionings for plasma facing walls of large
 helical device. Hino, T.; Ohuchi, T.; Hashiba, M.; Yamauchi, Y.;
 Hirohata, Y.; Inoue, N.; Sagara, A.; Noda, N.; Motojima, O.
 (Department of Nuclear Engineering, Hokkaido University, Sapporo,
 060-8628, Japan). Journal of Nuclear Materials, 290-293, 1176-1179
 (English) 2001. CODEN: JNUMAM. ISSN: 0022-3115.
 Publisher: Elsevier Science B.V..

AΒ The first and second exptl. campaigns in large helical device (LHD) were carried out from Mar. to Dec. 1998. Before the each campaign, the material probes were placed at the inner wall of vacuum vessel along the poloidal direction. After each campaign, change of surface morphol., impurity depth profile and gas desorption of the samples were examd. In the first exptl. campaign, the ECR discharge cleanings were employed. After the first exptl. campaign, the surface was modified by the deposition of sub-micron particles, and the concns. such as oxygen and carbon were still high. Impurity gas desorption was also large in the sample at the port. In the second exptl. campaign, the glow discharge cleanings were employed and the no. of main discharge shots increased. After the second exptl. campaign, no significant deposition took place except for the position close to the divertor leg, and the oxygen impurity level was reduced. In every sample, the helium was retained by the helium glow discharge. addn., the amt. of gas desorption was considerably reduced even in the sample at the port. In the second exptl. campaign, the wall conditionings largely progressed by using the glow discharge cleanings and the increase of main discharge shots with a high heating power.

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME)

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H<sub>2</sub>O
     7782-44-7 HCA
RN
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
CC
     71-2 (Nuclear Technology)
     Section cross-reference(s): 55
IT
     Desorption
     Electron cyclotron resonance
     Erosion (wear)
     Fusion reactor divertors
     Fusion reactor plasmas
     Impurities
        (conditionings for plasma facing walls of large helical
        device)
IT
     Fusion reactors
        (heliotron; conditionings for plasma facing walls of
        large helical device)
IT
     11134-23-9
        (conditionings for plasma facing walls of large helical
IT
     74-82-8, Methane, processes
                                   124-38-9, Carbon dioxide, processes
     630-08-0, Carbon monoxide, processes 1333-74-0, Hydrogen,
                 7440-59-7, Helium, processes 7732-18-5, Water,
     processes
     processes 7782-44-7, Oxygen, processes
        (conditionings for plasma facing walls of large helical
        device)
L66 ANSWER 12 OF 32 HCA COPYRIGHT 2006 ACS on STN
134:74032 Polymer electrolyte membrane with integrated
     plasma-polymerized ion-conducting barrier layer. Mueller,
     Joerg; Guth, Thomas; Mex, Laurent (Germany). Ger. Offen. DE
     19914571 A1 20010104, 4 pp.
                                   (German).
                                              CODEN: GWXXBX.
     APPLICATION: DE 1999-19914571 19990331.
AΒ
     The permeability of fuel cells with polymer electrolyte
     membranes to fuels (esp. methanol), is diminished sharply
     (vs. the state of the art) by coating the membrane
     surfaces with highly crosslinked plasma-polymd.
     ion-conducting layers.
IT
     7732-18-5, Water, processes
        (polymer electrolyte membrane with integrated
        plasma-polymd. ion-conducting barrier layer)
     7732-18-5 HCA
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RN

CN Water (8CI, 9CI) (CA INDEX NAME) H<sub>2</sub>O IT 1333-74-0, Hydrogen, uses 7782-44-7, Oxygen, uses (polymer electrolyte membrane with integrated plasma-polymd. ion-conducting barrier layer) RN1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-- H RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IC ICM H01M008-02 ICS H01M008-10; C08F002-46 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38 ST fuel cell polymer membrane fuel permeability decrease Polymerization IT (plasma; polymer electrolyte membrane with integrated plasma-polymd. ion-conducting barrier layer) IT Fuel cells Permeability Polymer electrolytes (polymer electrolyte membrane with integrated plasma-polymd. ion-conducting barrier layer) 630-08-0, Carbon monoxide, processes 7732-18-5, Water, IT processes (polymer electrolyte membrane with integrated plasma-polymd. ion-conducting barrier layer) 67-56-1, Methanol, uses 1333-74-0, Hydrogen, uses IT 7782-44-7, Oxygen, uses (polymer electrolyte membrane with integrated plasma-polymd. ion-conducting barrier layer) ANSWER 13 OF 32 HCA COPYRIGHT 2006 ACS on STN 133:307286 Biosensor using plasma-polymerized membrane Muguruma, Hitoshi; Hiratsuka, Akinori; Karube, Masao (Sentan Kaqaku Gijutsu Incubation Center K. K., Japan). Jpn. Kokai Tokkyo Koho JP 2000298111 A2 20001024, 11 pp. (Japanese).

CODEN: JKXXAF. APPLICATION: JP 1999-107691 19990415.

AB A highly functional biosensor with a novel structure is conveniently constructed using a plasma-polymd. membrane. The biosensor is constituted with a plasma-polymd. membrane contg. functional groups, a catalytically active substance (e.g., enzyme) immobilized on the plasma-polymd. membrane using a crosslinking reagent, and a metal electrode pattern in contact with a sample through the plasma -polymd. membrane. The influence by interfering compds. is eliminated due to the hydrogen peroxide-selective permeability of the membrane. The sensor can be applied in a wide range of areas in combination with micromachine technique. A diagram describing the sensor assembly is given. IT 1333-74-0, Hydrogen, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses (biosensor using plasma-polymd. membrane) RN1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H<sub>2</sub>O RN7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN 0 = 0IC ICM G01N027-327 CC 9-1 (Biochemical Methods) ST biosensor plasma polymn membrane enzyme electrode IT Amide group Amino group Carbonyl group Carboxyl group Crosslinking agents Enzyme electrodes Epoxy group Formyl group Functional groups Glucose sensors

Hydroxyl group

ΙT

IT

ΙT

IT

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Immobilization, biochemical
       Membrane electrodes
       Membranes, nonbiological
     Micromachines
     Permeability
     Sulfhydryl group
        (biosensor using plasma-polymd. membrane)
     Enzymes, uses
        (biosensor using plasma-polymd. membrane)
     Metals, uses
        (biosensor using plasma-polymd. membrane)
     Halogens
        (biosensor using plasma-polymd. membrane)
    Monomers
        (biosensor using plasma-polymd. membrane)
IT.
    Noble gases, uses
        (biosensor using plasma-polymd. membrane)
     Sensors
        (electrochem.; biosensor using plasma-polymd.
       membrane)
     Functional groups
        (imino group; biosensor using plasma-polymd.
       membrane)
     Functional groups
        (isocyanato group; biosensor using plasma-polymd.
       membrane)
     Polymerization
        (plasma; biosensor using plasma-polymd.
       membrane)
     Functional groups
        (vinyl group; biosensor using plasma-polymd.
       membrane)
     7722-84-1, Hydrogen peroxide, analysis
        (biosensor using plasma-polymd. membrane)
     9000-88-8, D-Amino acid oxidase 9000-89-9, L-Amino acid oxidase
    9001-37-0, Glucose oxidase 9001-46-1, Glutamate dehydrogenase
     9001-96-1, Pyruvate oxidase 9028-14-2, Glycerol dehydrogenase
     9028-53-9, Glucose dehydrogenase 9028-67-5, Choline oxidase
     9028-76-6, Cholesterol oxidase 9028-79-9, Galactose oxidase
     9028-86-8, Aldehyde dehydrogenase 9031-72-5, Alcohol dehydrogenase
     9035-73-8, Oxidase
                         9035-82-9, Dehydrogenase
                                                     9059-11-4, Amine
    oxidase
              67775-34-2, Cholesterol dehydrogenase
                                                       135622-84-3,
    Fructose dehydrogenase 220983-94-8, Sorbitol dehydrogenase
        (biosensor using plasma-polymd. membrane)
    50-81-7, Ascorbic acid, analysis
                                        51-61-6, Dopamine, analysis
    57-13-6, Urea, analysis
                               103-90-2
        (biosensor using plasma-polymd. membrane)
    7440-06-4, Platinum, uses
```

```
(biosensor using plasma-polymd. membrane)
ΙT
     302-01-2, Hydrazine, uses 1333-74-0, Hydrogen, uses
     7664-41-7, Ammonia, uses 7727-37-9, Nitrogen, uses
     7732-18-5, Water, uses 7782-44-7, Oxygen, uses
     7783-06-4, Hydrogen sulfide, uses 13465-07-1, Hydrogen disulfide
        (biosensor using plasma-polymd. membrane)
IT
     75-05-8, Acetonitrile, reactions
                                        107-46-0, Hexamethyldisiloxane
     111-30-8, Glutaraldehyde
        (biosensor using plasma-polymd. membrane)
     ANSWER 14 OF 32 HCA COPYRIGHT 2006 ACS on STN
L66
132:209793 Heat treatment of solid carbonaceous material in
     plasma reactors. Swanepoel, Jacobus; Lombaard, Ruan;
     Mast-Ingle, Julian Charles (S. Afr.). PCT Int. Appl. WO 2000013785
     A1 20000316, 34 pp. DESIGNATED STATES: W: AE, AL, AM,
     AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM,
     EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG,
     KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO,
     NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG,
     US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT,
     BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR,
     IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG.
                                                          (English).
                     APPLICATION: WO 1999-IB1477 19990826.
     CODEN: PIXXD2.
                                                            PRIORITY: ZA
     1998-8028 19980902.
AB
     An elec. arc plasma reactor is used
     for treating solid carbonaceous material by heating the material to
     ≥1800°C using a non-transfer arc generated
     plasma flame. Components in the carbonaceous material are
     gasified and removed from the residual solid material as a hot gas,
     where the residual solid material may be recovered as a product. A
     vertical shaft non-transfer arc plasma reactor is used
     which comprises an upper preheating zone, an intermediate reaction
     zone in which ≥1 non-transfer arc plasma generator
     or reactor is located, and a lower cooling zone.
IT
     1333-74-0, Hydrogen, processes 7782-44-7, Oxygen,
     processes
        (heat treatment of solid carbonaceous material in plasma
        reactors)
RN
     1333-74-0
               HCA
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-- H
RN
     7782-44-7 HCA
```

CN

Oxygen (8CI, 9CI) (CA INDEX NAME)

0==0

Tungsten, uses

IT 7732-18-5, Water, processes (vapor; heat treatment of solid carbonaceous material in **plasma** reactors) RN 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H<sub>2</sub>O IC ICM B01J019-08 ICS B01J008-18; B01J008-12; B01J008-08; C10B019-00; C10J003-18 CC 48-8 (Unit Operations and Processes) Section cross-reference(s): 49, 55, 56, 57, 76 carbonaceous material heat treatment plasma reactor ST Carburizing IT Ceramics Graphitization Heat treatment Linings (refractory) Reduction Thermal decomposition (heat treatment of solid carbonaceous material in plasma reactors) Carbides IT Metals, processes (heat treatment of solid carbonaceous material in plasma reactors) IT Carbonaceous materials (technological products) (heat treatment of solid carbonaceous material in plasma reactors) IT Coke (low-sulfur, low-ash; heat treatment of solid carbonaceous material in **plasma** reactors) IT (pitch; heat treatment of solid carbonaceous material in plasma reactors) IT Reactors (plasma; heat treatment of solid carbonaceous material in **plasma** reactors) IT 7440-44-0P, Carbon, processes 7782-42-5P, Graphite, processes (heat treatment of solid carbonaceous material in plasma reactors) IT 1314-23-4, Zirconia, uses 7439-98-7, Molybdenum, uses 7440-33-7, (heat treatment of solid carbonaceous material in **plasma** reactors)

- TT 74-90-8P, Hydrogen cyanide, processes 75-73-0P, Carbon tetrafluoride 76-16-4P 116-14-3P, Tetrafluoroethylene, processes 116-15-4P, Hexafluoropropylene 409-21-2P, Silicon carbide, processes 11114-46-8P, Ferrochromium (heat treatment of solid carbonaceous material in plasma reactors)
- IT 630-08-0, Carbon monoxide, processes 1333-74-0, Hydrogen, processes 7440-01-9, Neon, processes 7440-37-1, Argon, processes 7440-59-7, Helium, processes 7631-86-9, Silica, processes 7727-37-9, Nitrogen, processes 7782-41-4, Fluorine, processes 7782-44-7, Oxygen, processes 7782-50-5, Chlorine, processes 12068-77-8, Iron chromite (FeCr204) (heat treatment of solid carbonaceous material in plasma reactors)
- L66 ANSWER 15 OF 32 HCA COPYRIGHT 2006 ACS on STN
  131:345219 Applications of the cavity ring-down technique to a
   large-area rf-plasma reactor. Grangeon, F.;
   Monard, C.; Dorier, J.-L.; Howling, A. A.; Hollenstein, Ch.;
   Romanini, D.; Sadeghi, N. (Ecole Polytechnique Federale de Lausanne,
   Centre de Recherches en Physique des Plasmas, Lausanne, 1015,
   Switz.). Plasma Sources Science & Technology, 8(3), 448-456
   (English) 1999. CODEN: PSTEEU. ISSN: 0963-0252.
   Publisher: Institute of Physics Publishing.
- The cavity ring-down technique is applied to an industrial-scale radiofrequency (rf) plasma reactor for the measurement of the d. and spatial profile of neg. ions in pure 0 and H radiofrequency plasmas, and for the detection of nanometric particles in Ar-silane plasmas. The real-time observation of powder formation is feasible by the cavity ring-down technique. An obsd. plasma-induced spurious drift of the ring-down time is also studied and related to H2O desorption from the reactor walls and electrodes which is re-adsorbed on the mirror surfaces.
- IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties

(applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d.)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

RN · 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IT 7732-18-5, Water, processes (desorption; applications of cavity ring-down technique to large-area RF-plasma reactor with) RN 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) CN H20 CC 76-11 (Electric Phenomena) cavity ring plasma reactor; anion formation plasma ST reactor oxygen hydrogen; silane argon plasma reactor anion IT Mirrors (applications of cavity ring-down technique to large-area RF-plasma reactor) IT Anions Electric discharge devices (applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d.) IT Nanoparticles (applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d. and) Powders IT (applications of cavity ring-down technique to large-area RF-plasma reactor in deposition of) IT Degassing (applications of cavity ring-down technique to large-area RF-plasma reactor with) IT Photoionization (in applications of cavity ring-down technique to large-area **RF-plasma** reactor) IT Electrodes (plasma; applications of cavity ring-down technique to large-area RF-plasma reactor) IT Reactors (plasma; applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d.) 12184-88-2, Hydride 14337-01-0, Oxygen(1-), properties IT (applications of cavity ring-down technique to large-area

RF-plasma reactor for measurement of anion d.)

IT 7440-37-1, Argon, processes 7803-62-5, Silane, processes (applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d.)

IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties

(applications of cavity ring-down technique to large-area RF-plasma reactor for measurement of anion d.)

IT 7732-18-5, Water, processes

(desorption; applications of cavity ring-down technique to large-area RF-plasma reactor with)

- L66 ANSWER 16 OF 32 HCA COPYRIGHT 2006 ACS on STN
- 131:93862 Boron-doped CVD diamond as an electrode material for cyclic voltammetry in various electrolyte solutions for electroanalytical applications. Ramesham, R.; Rose, M. F. (Space Power Institute, Auburn University, Auburn, AL, 36849-5320, USA). High Temperature and Materials Science, Volume Date 1997, 38(1), 1-12 (English) 1999. CODEN: HTMSFP. ISSN: 1080-1278. Publisher: Humana Press Inc..
- AB Boron-doped polycryst. diamond films were grown over a molybdenum substrate by a microwave plasma CVD process using a methane and hydrogen gas mixt. at a pressure of 35.7 torr. Boron doping of diamond was achieved in situ by using a solid boron source while growing diamond in the CVD process. Cyclic voltammetry was used to evaluate the boron-doped diamond-coated molybdenum electrode material in 0.5M NaCl, 0.5M HNO3, 0.5M HCl, 0.5M KOH, 1M KNO3, and 1M Na2SO4 solns. The authors obsd. a negligible background current for diamond by cyclic voltammetry in the various solns. over a wide potential range. Therefore, diamond will certainly have a use as an electrode material in electroanal. applications. The authors have studied the redox kinetic behavior of Fe(CN)63- + e- <=> Fe(CN)64- at the boron-doped diamond electrode in 1M KNO3 supporting electrolyte soln. and cor. the voltammetric response for IR compensation. The heterogeneous electron transfer rate const. was detd. using the exptl. data and COOL algorithm. rate const. is approx. 2 + 10-4 cm/s, and the kinetics were considered to be sluggish at the diamond electrode/soln. interface.

IT 7732-18-5, Water, processes

(electrolysis; on boron-doped diamond electrode)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties

```
(evolution on boron-doped diamond electrode in aq. solns.)
RN
     1333-74-0
CN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
H-H
RN
     7782-44-7 HCA
CN
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
CC
     72-2 (Electrochemistry)
     Section cross-reference(s): 67, 75, 79
IT
     Vapor deposition process
        (plasma; of boron-doped diamond for electrode)
IT
     7732-18-5, Water, processes
        (electrolysis; on boron-doped diamond electrode)
IT
     1333-74-0, Hydrogen, properties 7782-44-7, Oxygen,
     properties
        (evolution on boron-doped diamond electrode in aq. solns.)
IT
     74-82-8, Methane, reactions
        (in microwave plasma CVD of boron-doped
        diamond for electrode)
     ANSWER 17 OF 32 HCA COPYRIGHT 2006 ACS on STN
131:89695 Process and system for separation and recovery of
     perfluorocompound gases. Li, Yao-En; Paganessi, Joseph E.;
     Vassallo, David; Fleming, Gregory K. (L'Air liquide S. A, Fr.).
                                                                       PCT
     Int. Appl. WO 9936159 A1 19990722, 105 pp. DESIGNATED
                AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU,
     STATES: W:
     CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE,
     KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
     NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA,
     UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT,
     BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR,
     IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG.
                                                          (English).
     CODEN: PIXXD2.
                     APPLICATION: WO 1998-EP2272 19980416. PRIORITY: JP
     1998-7021 19980116.
AB
     Processes and systems to recover at least one perfluorocompound gas
     from a gas mixt. are provided. In one embodiment the inventive
     process comprises providing a gas mixt. comprising at least one
     perfluorocompound gas and at least one carrier gas, the gas mixt.
    being at a predetd. pressure; providing at least one size selective
    membrane having a feed side and a permeate side; contacting
     the feed side of the at least one membrane with the gas
     mixt.; withdrawing from the feed side of the membrane as a
```

IT

RN

CN

H-H

RN

CN

H<sub>2</sub>O

RN CN

IC

CC

IT

IT

IT

gases)

non-permeate stream at a pressure which is substantially equal to the predetd. pressure a concd. gas mixt. comprising essentially the at least one perfluorocompound gas; and withdrawing from the permeate side of the membrane as a permeate stream a depleted gas mixt. comprising essentially the at least one carrier gas. 1333-74-0, Hydrogen, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses (process and system for sepn. and recovery of perfluorocompound 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) 7732-18-5 HCA Water (8CI, 9CI) (CA INDEX NAME) 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0ICM B01D053-22 48-1 (Unit Operations and Processes) Section cross-reference(s): 59, 76 Air purification (membrane sepn.; process and system for sepn. and recovery of perfluorocompound gases) Decomposition (plasma; process and system for sepn. and recovery of perfluorocompound gases) Adsorption Gases Heat exchangers Membranes, nonbiological Scrubbers Scrubbing Semiconductor device fabrication Separation Thermal decomposition (process and system for sepn. and recovery of perfluorocompound

- 17 124-38-9, Carbon dioxide, uses 630-08-0, Carbon monoxide, uses 1333-74-0, Hydrogen, uses 7439-90-9, Krypton, uses 7440-01-9, Neon, uses 7440-37-1, Argon, uses 7440-59-7, Helium, uses 7440-63-3, Xenon, uses 7727-37-9, Nitrogen, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses (process and system for sepn. and recovery of perfluorocompound gases)
- ANSWER 18 OF 32 HCA COPYRIGHT 2006 ACS on STN L66 130:30543 Preliminary study of ferri-ferrocyanide redox kinetics at the submillimeter boron-doped CVD diamond electrode. Ramesham, Rajeshuni (Space Power Institute, 231 Leach Center, Auburn University, Auburn, AL, 36849-5320, USA). Sensors and Materials, 10(5), 275-286 (English) 1998. CODEN: SENMER. ISSN: 0914-4935. Publisher: Scientific Publishing Division of MYU K.K.. ABBoron-doped CVD diamond was grown on mech. damaged and cleaned molybdenum substrate using the microwave plasma CVD process. The gases used to grow the CVD diamond were methane and hydrogen. Boron doping of the diamond was achieved in situ using a solid disk source of B2O3. The submillimeter boron-doped CVD diamond electrode was evaluated for background current response in 0.5M NaCl by cyclic voltammetry. Photolithog. was used to pattern the CVD diamond to fabricate a submillimeter electrode for cyclic voltammetry studies. Preliminary results on the voltammetric response of the fabricated (0.003 cm2) CVD diamond electrode are reported. The authors obsd. negligible background current for submillimeter boron-doped CVD diamond electrode over a wide potential range. Decompn. of water occurs electrochem., evolving O2 during pos. (anodic) polarization and H2 during neg. (cathodic) polarization. SEM was used to study the morphol. of the diamond electrode before and after electrochem. characterization; the authors conclude qual. that the morphol. was unchanged. The authors have studied the redox kinetics of ferri-ferrocyanide (Fe(CN)63- + e- .dblarw. Fe(CN)64-) in 0.5M NaCl soln. at various scan rates (1 mV/s to 20,000 mV/s). The reaction is reversible at the diamond electrode, but the redox kinetics are slow. The results of cyclic voltammetry using submillimeter boron-doped CVD diamond electrode reveal a change in shape of the voltammetric response curve from sigmoidal to peak-shaped as a function of scan rate, reflecting the
- IT 7732-18-5, Water, processes

(electrolysis; hydrogen and oxygen evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped CVD diamond electrode)

- RN 7732-18-5 HCA
- CN Water (8CI, 9CI) (CA INDEX NAME)

change in the diffusion layer thickness.

H<sub>2</sub>O IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties (evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped CVD diamond electrode) 1333-74-0 HCA RNHydrogen (8CI, 9CI) (CA INDEX NAME) CNH-HRN7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IT7732-18-5, Water, properties (hydrogen and oxygen evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped CVD diamond electrode) 7732-18-5 HCA RN Water (8CI, 9CI) (CA INDEX NAME) CN H20 CC 72-2 (Electrochemistry) Section cross-reference(s): 67, 75 cyanoferrate electroredox kinetics boron doped diamond; electrode ST boron doped diamond cyanoferrate redox; plasma CVD boron doped diamond electrode Vapor deposition process IT (plasma; of boron-doped diamond for electrode for ferri-ferrocyanide redox kinetics) IT 7732-18-5, Water, processes (electrolysis; hydrogen and oxygen evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped CVD diamond electrode) 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, IT properties (evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped

CVD diamond electrode)

IT 7732-18-5, Water, properties (hydrogen and oxygen evolution in water electrolysis on boron-doped diamond: ferri-ferrocyanide redox kinetics at submillimeter boron-doped CVD diamond electrode)

L66 ANSWER 19 OF 32 HCA COPYRIGHT 2006 ACS on STN 129:253583 CVD and etching process using RF induction plasma and apparatus therefor. Okumura, Tomohiro; Tanabe, Hiroshi; Sawada, Kazuyuki; Arai, Koji; Suzuki, Naoki (Matsushita Electric Industrial Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 10261630 A2 **19980929** Heisei, 9 pp. (Japanese). JKXXAF. APPLICATION: JP 1997-65984 19970319.

AB The process is carried out in a condition where a source (or etching) gas (G1) and a gas (G2) which inhibits deposition of G1 on a dielec. members are flown in a vacuum chamber. The coil, being applied with RF power to generate a plasma, is arranged along to the dielec. member. Also claimed is the app. for the process. The undesirable adhesion of CVD source or etching gas on the dielec. member is prevented to improve the processing reproducibility.

ΙT 7732-18-5, Water, uses 7782-44-7, Oxygen, uses (deposition-inhibiting gas; plasma processing method and its app. inhibiting film deposition on dielec. substance surface)

7732-18-5 HCA RN

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0===0

1333-74-0, Hydrogen, uses 1333-74-0D, Hydrogen, IT halide, uses (etching gas; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) RN 1333-74-0 HCA

Hydrogen (8CI, 9CI) (CA INDEX NAME) CN

H-H

RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-HIC ICM H01L021-31 ICS C23C016-52; C23F004-00; H01L021-205; H01L021-285; H01L021-3065 CC 76-14 (Electric Phenomena) ST CVD etching app RF induction plasma; dielec coil member source adhesion prevention; argon deposition preventing gas plasma app IT Electric coils (plasma processing method and its app. inhibiting film deposition on dielec. substance surface) ITVapor deposition apparatus (plasma; CVD and etching process using RF induction plasma and app. therefor) IT Etching apparatus (plasma; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) IT 7440-59-7, Helium, uses 7440-37-1, Argon, uses 7727-37-9. Nitrogen, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses 10024-97-2, Nitrous oxide, uses (deposition-inhibiting gas; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) IT 75-10-5, Difluoromethane 75-46-7, Trifluoromethane 75-73-0, Perfluoromethane 1333-74-0, Hydrogen, uses 1333-74-0D, Hydrogen, halide, uses Perfluorobutene 51311-17-2, Carbon fluoride (etching gas; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) IT 7631-86-9P, Silica, processes (plasma processing method and its app. inhibiting film deposition on dielec. substance surface) 7440-32-6, Titanium, processes 7440-33-7, Tungsten, processes IT (source contg.; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) IT 78-10-4, Tetraethoxysilane (source; plasma processing method and its app. inhibiting film deposition on dielec. substance surface) L66 ANSWER 20 OF 32 HCA COPYRIGHT 2006 ACS on STN 129:167244 A study on the origin of nonfaradaic behavior of anodic contact glow discharge electrolysis, The relationship between power dissipated in glow discharges and nonfaradaic yields. Sengupta, Susanta K.;

Singh, Rajeshwar; Srivastava, Ashok K. (Department of Chemistry, Faculty of Science, Banaras Hindu University, Varanasi, 221005,

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India). Journal of the Electrochemical Society, 145(7), 2209-2213
     (English) 1998. CODEN: JESOAN.
                                      ISSN: 0013-4651.
     Publisher: Electrochemical Society.
     Chem. effects of contact glow discharge
AB
     electrolysis (CGDE) at an electrode where a plasma is
     sustained by d.c. glow discharges between the
     electrode and the surrounding electrolyte, are remarkably
     nonfaradaic. A crit. anal. of the chem. results of anodic CGDE at
     varying voltages, currents, power supplies, and pHs clearly shows
     that nonfaradaic yields originate in two sep. reaction zones: the
     plasma around the anode and the liq. anolyte near the
     plasma-anolyte interface. The yields from the former zone
     appear from 250 V onward (the beginning of the onset of partial
     glow discharge) and vary linearly with the power
     dissipated in the glow discharge. The yields
     from the latter zone appear from 410 V onward (the beginning of the
     full glow discharge) and are independent of the
     power dissipated in the glow discharge.
     relative contribution of the two zones to the total nonfaradaic
     yields is dependent on the voltage applied: the plasma
     zone having a share of 100% up to 400 V, followed by 20% up to 450
     V, and thereafter rising steadily to 57% at 500 V.
IT
     7732-18-5, Water, processes
        (electrolysis; anodic contact glow discharge
        electrolysis)
RN
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
CN
H<sub>2</sub>O
IT
     1333-74-0, Hydrogen, formation (nonpreparative)
     7782-44-7, Oxygen, formation (nonpreparative)
        (formation in anodic contact glow discharge
        electrolysis in phosphate buffered solns.)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
o = o
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CC

72-2 (Electrochemistry)

Section cross-reference(s): 76
ST nonfaradaic anodic contact glow discharge electrolysis; power dissipation glow discharge

nonfaradaic yield

IT Plasma

(in anodic contact glow discharge electrolysis)

IT Electrolysis

Glow discharge

Power

(origin of nonfaradaic behavior of anodic contact **glow discharge** electrolysis: relationship between power dissipated in **glow discharges** and nonfaradaic yields)

IT 1310-58-3, Potassium hydroxide, uses (anodic contact **glow discharge** electrolysis of phosphate buffered soln. of)

1333-74-0, Hydrogen, formation (nonpreparative) 7722-84-1, Hydrogen peroxide, formation (nonpreparative) 7782-44-7, Oxygen, formation (nonpreparative) (formation in anodic contact glow discharge electrolysis in phosphate buffered solns.)

L66 ANSWER 21 OF 32 HCA COPYRIGHT 2006 ACS on STN

128:194377 Recovery of noble gases from plasma display panel sealing furnace. Jain, Ravi; Whitlock, Walter H. (BOC Group, Inc., USA). Eur. Pat. Appl. EP 826629 A2 19980304, 14 pp.

DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO. (English). CODEN: EPXXDW. APPLICATION: EP 1997-305948 19970805. PRIORITY: US 1996-703711 19960827; US 1997-884421 19970627.

AB A noble gas selected from xenon, neon, krypton or their mixts. is recovered from a process chamber by purging the chamber with one of several gases and then sepg. the noble gas from the purge gas by one of several techniques. The sepd. noble gas is further purified to remove trace impurities and is then recycled for reuse in the process chamber. The purge gas may be recycled or disposed of, depending upon the particular purge gas used and its purity. In a specific embodiment, hydrogen is used as the purge gas and it is sepd. from the noble gas by high temp. membrane sepn. using a palladium membrane. The hydrogen, obtained in high purity, and the noble gas are each recycled for reuse in the

process chamber. IT 1333-74-0, Hydrogen, uses 7782-44-7, Oxygen, uses (purging gas; in recovery of noble gases from plasma display panel sealing furnace) RN 1333-74-0 HCA CN Hydrogen (8CI, 9CI) (CA INDEX NAME) H-HRN7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0ICM C01B023-00 IC ICS B01D053-00; B01D053-02; B01D053-22 49-1 (Industrial Inorganic Chemicals) CC ST noble gas recovery hydrogen purging; neon recovery hydrogen purging; xenon recovery hydrogen purging; krypton recovery hydrogen purging; plasma display panel sealing furnace Hydrocarbons, uses IT (C4-6 purging gas; recovery of noble gases from plasma display panel sealing furnace) IT Steam (purging gas; recovery of noble gases from plasma display panel sealing furnace) Recycling IT (recovery of noble gases from plasma display panel sealing furnace) IT Noble gases, preparation (recovery of noble gases from plasma display panel sealing furnace) IT 74-85-1, Ethylene, uses 115-07-1, Propylene, uses Carbon dioxide, uses 630-08-0, Carbon monoxide, uses 1333-74-0, Hydrogen, uses 7664-41-7, Ammonia, uses 7782-44-7, Oxygen, uses (purging gas; in recovery of noble gases from plasma display panel sealing furnace) 7439-90-9P, Krypton, preparation IT 7440-01-9P, Neon, preparation 7440-63-3P, Xenon, preparation (recovery of noble gases from plasma display panel sealing furnace) ANSWER 22 OF 32 HCA COPYRIGHT 2006 ACS on STN

126:311125 Discharge methods and electrodes for generating plasmas at one atmosphere of pressure, and materials treated therewith.

AΒ

IT

RN

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H-H

RN

CN

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Spence, Paul D. (University of Tennessee Research Corporation, USA).
       PCT Int. Appl. WO 9713266 A2 19970410, 63 pp.
     DESIGNATED STATES: W: AU, BR, CA, CN, DE, DE, IL, JP, KR, RU; RW:
     AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE.
     (English). CODEN: PIXXD2. APPLICATION: WO 1996-US20919 19960619.
     PRIORITY: US 1995-492193 19950619.
     Two methods and corresponding electrode designs are provided for the
     generation of a plasma at .apprx.1 atm. Using these
     methods, various webs, films, and 3-dimensional objects, esp. of
     polymers, are beneficially treated (e.q., to improve wettability or
     printability) in a reduced amt. of time. A 1st method uses a
     repetitive, asym. voltage pulse to generate a plasma
     discharge between 2 electrodes. An asym. voltage pulse is used to
     generate a discharge in which a substrate can be exposed
     predominantly to either pos. or neg. plasma species,
     depending on the voltage polarity used. A 2nd method uses the gap
     capacitance of an electrode pair and an external inductor in shunt
     to form a resonant LC circuit. The circuit is driven by a
     high-power radio-frequency source operating at
     1-30 MHz to generate a uniform discharge between the electrode pair.
     Both methods have temp.-controlled discharge surfaces with supply
     gas temp., humidity, and flow rate control. The gas flow is
     typically sufficient to cause a turbulent flow field in the
     discharge region where materials are treated. Electrode pairs
     implement these methods and include a metal-faced electrode and a
     dielec.-covered electrode, 1 or both of which have holes extending
     through the electrode face for supply gas flow. The 2nd of the
     above-described methods will also operate with paired, metal-faced
     electrodes, but under more restricted operating conditions.
    1333-74-0, Hydrogen, processes 7782-44-7, Oxygen,
    processes
        (app. for generating plasmas for surface treatment of)
     1333-74-0 HCA
    Hydrogen (8CI, 9CI) (CA INDEX NAME)
     7782-44-7 HCA
    Oxygen (8CI, 9CI) (CA INDEX NAME)
0==0
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ΙT 7732-18-5, Water, processes (vapor; app. for generating plasmas for surface treatment of) RN7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

- IC ICM H01J
- CC 76-11 (Electric Phenomena)
   Section cross-reference(s): 38
- ST discharge electrode method plasma generation; polymer plasma treatment
- IT Electrodes

## Plasma

(discharge methods and electrodes for generating plasmas for surface treatment)

- IT 124-38-9, Carbon dioxide, processes 1333-74-0, Hydrogen, processes 7440-37-1, Argon, processes 7440-59-7, Helium, processes 7664-41-7, Ammonia, processes 7727-37-9, Nitrogen, processes 7782-44-7, Oxygen, processes 9002-88-4, Polyethylene 9003-07-0, Polypropylene 9003-53-6, Polystyrene 9010-79-1, Ethylene-propylene copolymer 24968-12-5, Polybutylene terephthalate 25038-59-9, processes 26062-94-2, Polybutylene terephthalate
  - (app. for generating plasmas for surface treatment of)
- IT 7732-18-5, Water, processes

(vapor; app. for generating plasmas for surface treatment of)

- L66 ANSWER 23 OF 32 HCA COPYRIGHT 2006 ACS on STN
- 125:97909 The wall conditioning techniques in RFX. Sonato, P.; Baker, W. R.; Fiorentin, P.; Marchiori, G.; Zaccaria, P.; Zollino, G.; Antoni, V.; Bagatin, M.; Desideri, D.; et al. (Gruppo di Padova per ricerche sulla Fusione, EURATOM-ENEA-CNR-Universita di Padova, Padua, 35127, Italy). Vacuum, 47(6-8, Proceedings of the 13th International Vacuum Congress and the 9th International Conference on Solid Surfaces, 1995), 977-980 (English) 1996. CODEN: VACUAV. ISSN: 0042-207X. Publisher: Elsevier.
- The vacuum of RFX has a total vol. of 10 m3 and a surface exposed to the vacuum of 40 m2. It is completely covered with 2016 graphite tiles. The following wall conditioning techniques are applied: baking up to 350°, GDC and PDC in H2 and He. Moreover, in order to reduce Zeff in the plasma discharge, a GDC assisted deposition of a film contg. boron has been carried out. The effectiveness is assessed by means of mass spectrometers and by silicon samples exposed to the processes. In this paper, after a brief description of the wall conditioning techniques, the main results are reported and compared.
- IT **1333-74-0**, Hydrogen, uses

(glow and pulse discharge cleaning as wall

conditioning techniques for the RFX reversed field pinch) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-HIT 7732-18-5, Water, processes 7782-44-7, Oxygen, processes (wall conditioning techniques used in the RFX reversed field pinch in relation to) 7732-18-5 HCA RNCN Water (8CI, 9CI) (CA INDEX NAME) H<sub>2</sub>O RN 7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN0 = 071-2 (Nuclear Technology) CC reversed field pinch wall conditioning RFX; fusion reactor wall ST conditioning RFX; pulse discharge cleaning wall conditioning pinch; baking wall conditioning reversed field pinch; glow discharge cleaning wall conditioning pinch; boronization wall conditioning reversed field pinch IT Cleaning (glow discharge, wall conditioning techniques for the RFX reversed field pinch) 1333-74-0, Hydrogen, uses 7440-59-7, Helium, uses IT (glow and pulse discharge cleaning as wall conditioning techniques for the RFX reversed field pinch) IT 124-38-9, Carbon dioxide, processes 630-08-0, Carbon monoxide, processes 7732-18-5, Water, processes 7782-44-7, Oxygen, processes (wall conditioning techniques used in the RFX reversed field pinch in relation to) L66 ANSWER 24 OF 32 HCA COPYRIGHT 2006 ACS on STN 124:71363 Process-induced damage to SRAM poly-load resistance during photoresist ashing in H2O plasma. Chang, Kuang-Hui; Huang, Yuang-Chang; Lin, Ting-Hwang; Chang, Chaur-Rong (Taiwan Semiconductor Manufacturing Company, Hsin Chu, Taiwan). Proceedings of SPIE-The International Society for Optical Engineering, 2635, 276-83 (English) 1995. CODEN: PSISDG. ISSN: 0277-786X.

Process-induced damage to SRAM (static-RAM) poly-load resistance AB have been clearly obsd. during photoresist ashing in an H2O plasma which is commonly used as an effective corrosion-prevented treatment after metal etching in a chlorine rich environment. The resistance degraded to about three order in the H2O plasma than in the conventional O2 plasma. Different ratios of H2O to O2 plasma have been studied by using the well-designed pattern to understand the basic mechanisms of the problems which occurred in the photoresist ashing process. Exptl. results showed that the higher concn. of H2O, the more damage to the SRAM products, resulting in the lower resistance of poly-load. Different types of plasma sources ( microwave and inductively coupled plasmas) for H2Oplasma photoresist ashing process have also been investigated and found that the damage effects to SRAM poly-load resistance in the inductive-type plasma source (such as Transformer Coupled Plasma) is more severe than that in the microwave-type plasma at the similar operation conditions. This may because inductive-type plasma has higher degree of ionization which generated more hydrogen ions inside the H2O plasma. 1333-74-0, Hydrogen, processes IT. (plasma; process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma) RN1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-H7732-18-5, Water, processes 7782-44-7, IT Oxygen, processes (plasma; process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma) RN 7732-18-5 HCA CN Water (8CI, 9CI) (CA INDEX NAME) H20 RN7782-44-7 HCA Oxygen (8CI, 9CI) (CA INDEX NAME) CN 0 = 0

74-5 (Radiation Chemistry, Photochemistry, and Photographic and

CC

Other Reprographic Processes)

ST photoresist ashing water plasma SRAM

IT Resists

(photo-, process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma)

IT Memory devices

(random-access, static; process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma)

IT 1333-74-0, Hydrogen, processes

(plasma; process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma)

IT 75-73-0, Carbon tetrafluoride 7732-18-5, Water,

processes 7782-44-7, Oxygen, processes

(plasma; process-induced damage to SRAM poly-load resistance during photoresist ashing in water plasma)

- L66 ANSWER 25 OF 32 HCA COPYRIGHT 2006 ACS on STN
- 122:108344 Surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of the membrane. Zhang, Liangren; Jin, Daosen; Tisato, Kajiyama (Lanzhou Inst. Chem. Phys., Acad. Sin., Lanzhou, 730000, Peop. Rep. China). Gaofenzi Xuebao (6), 660-6 (Chinese) 1993. CODEN: GAXUE9. ISSN: 1000-3304.
- AB The surface modification of cis-1,4-polybutadiene membrane was studied using CF2Cl2, CF2ClCF2Cl, and CF4 plasma treatments. The plasma-treated composite membrane was investigated by SEM, measurement of static contact angle to water, and XPS. Gas permeability of the membrane was also measured.
- 7732-18-5, Water, properties
   (contact angle of water to cis-1,4-polybutadiene with surface
   modified by halocarbon plasmas and gas permeability of
   membrane)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties

(surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

0 = 0

CC 39-12 (Synthetic Elastomers and Natural Rubber)

IT Contact angle

(contact angle of water to cis-1,4-polybutadiene with surface modified by halocarbon plasmas and gas permeability of membrane)

IT Permeability and Permeation

(gas permeability of cis-1,4-polybutadiene composite membrane with surface modified by halocarbon plasmas)

IT Plasma

Surface

(surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

IT Rubber, butadiene, properties

(of cis-1,4-configuration, surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

IT 7732-18-5, Water, properties

(contact angle of water to cis-1,4-polybutadiene with surface modified by halocarbon plasmas and gas permeability of membrane)

IT 9003-17-2

(rubber, of cis-1,4-configuration, surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of **membrane**)

IT 124-38-9, Carbon dioxide, properties 630-08-0, Carbon monoxide, properties 1333-74-0, Hydrogen, properties 7727-37-9,

Nitrogen, properties 7782-44-7, Oxygen, properties

(surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

IT 9003-17-2

(surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

IT 75-73-0, Tetrafluoromethane 76-14-2, 1,2-Dichloro-1,1,2,2-tetrafluoroethane

(surface modification of cis-1,4-polybutadiene with halocarbon plasmas and gas permeability of membrane)

IT 75-71-8, Dichlorodifluoromethane

(surface modification of cis-1,4-polybutadiene with halocarbon

plasmas and gas permeability of the membrane)

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L66 ANSWER 26 OF 32 HCA COPYRIGHT 2006 ACS on STN
122:85486 Gas plasma-treated solid polymer electrolyte
     membrane for fuel cell and fuel cell including this
     membrane. Binder, Michael; Mammone, Robert J. (United
     States Dept. of the Army, USA). U.S. US 5372896 A 19941213
     , 3 pp. (English). CODEN: USXXAM. APPLICATION: US 1993-123652
     19930920.
     A perfluorosulfonate membrane is exposed for .apprx.8 min
AB
     to a gas plasma prior to inclusion in a fuel cell as a
     separator between a liq. or gaseous fuel and an oxidant.
IT
     1333-74-0, Hydrogen, uses 7732-18-5, Water, uses
     7782-44-7, Oxygen, uses
        (fuel-cell perfluorosulfonate electrolyte membrane gas
        plasma treated with)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
RN
     7732-18-5 HCA
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H20
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
o = o
     ICM H01M008-10
IC
INCL 429033000
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     Section cross-reference(s): 38
     fuel cell polymer electrolyte membrane; gas plasma
ST
     treatment electrolyte membrane; perfluorosulfonate
    membrane gas plasma treatment
IT
    Electric conductivity and conduction
        (ionic, of gas plasma-treated fuel-cell
       perfluorosulfonate electrolyte membranes)
IT
    Fuel cells
        (separators, gas plasma-treated perfluorosulfonate
       membrane for)
IT 75-73-0, Tetrafluoromethane 124-38-9, Carbon dioxide, uses
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IT

AB

IT

RNCN

H-H

RN

CN

H<sub>2</sub>O

RN

CN

IC

ICM B01D067-00

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1333-74-0, Hydrogen, uses
                               7440-59-7, Helium, uses
     7446-09-5, Sulfur dioxide, uses 7664-41-7, Ammonia, uses
     7732-18-5, Water, uses 7782-44-7, Oxygen, uses
     10024-97-2, Nitrogen oxide (N2O), uses 10028-15-6, Ozone, uses
        (fuel-cell perfluorosulfonate electrolyte membrane gas
        plasma treated with)
     66796-30-3, Nafion 117
        (gas plasma-treated fuel-cell electrolyte
        membrane of)
    ANSWER 27 OF 32 HCA COPYRIGHT 2006 ACS on STN
121:12419 Manufacture of semipermeable polymer membranes.
     Gadkaree, Kishor Purushottam; Hersch, Leroy S. (Corning, Inc., USA).
       Eur. Pat. Appl. EP 595201 A2 19940504, 10 pp.
     DESIGNATED STATES: R: DE, FR, GB. (English). CODEN: EPXXDW.
     APPLICATION: EP 1993-117068 19931021. PRIORITY: US 1992-968661
     The title microporous polymer membranes, useful as a sepn.
     membrane or the like, are prepd. by extruding a melt-blended
     mixt. of a high-temp. thermoplastic polymer and a leachable glass to
     form sheet and leaching the glass from the sheet to leave a
     microporous relict polymer network. The permeability of the sheet
     is improved by plasma-etching the surface of the sheet
     before or after leaching. Hydrophilic or hydrophobic porous
     membranes may be provided.
     1333-74-0, Hydrogen, uses 7732-18-5, Water, uses
     7782-44-7, Oxygen, uses
        (plasma, in manuf. of semipermeable microporous polymer
        membranes)
     1333-74-0 HCA
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
     7732-18-5 HCA
     Water (8CI, 9CI) (CA INDEX NAME)
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
0 = 0
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ICS B01D069-00

CC 47-2 (Apparatus and Plant Equipment)
 Section cross-reference(s): 38, 57, 61

- ST polymer membrane semipermeable manuf; glass leachable polymer membrane manuf; plasma etching polymer membrane manuf
- IT Polysulfones, uses

(membranes, semipermeable microporous, manuf. of, leachable glass and plasma etching in)

IT Sputtering

(etching, in manuf. of semipermeable microporous polymer membranes)

IT Glass, oxide

(phosphate, leachable, in manuf. of semipermeable microporous polymer membranes)

IT Polyethers, uses

(polyamide-, membranes, semipermeable microporous, manuf. of, leachable glass and plasma etching in)

IT Polyamides, uses

Polyketones

(polyether-, membranes, semipermeable microporous, manuf. of, leachable glass and plasma etching in)

IT Polyethers, uses

(polyketone-, membranes, semipermeable microporous, manuf. of, leachable glass and plasma etching in)

IT Membranes

(semipermeable, polymer, manuf. of microporous, leachable glass and plasma etching in)

IT Etching

(sputter, in manuf. of semipermeable microporous polymer membranes)

IT 1333-74-0, Hydrogen, uses 7664-41-7, Ammonia, uses 7732-18-5, Water, uses 7782-44-7, Oxygen, uses 10024-97-2, Nitrogen oxide (N2O), uses

(plasma, in manuf. of semipermeable microporous polymer membranes)

- L66 ANSWER 28 OF 32 HCA COPYRIGHT 2006 ACS on STN
- 113:60252 Low pressure plasma chemistry in the Institute of Chemistry, Academia Sinica. Chen, Guanwen (Inst. Chem., Acad. Sin., Beijing, Peop. Rep. China). Proceedings of Japanese Symposium on Plasma Chemistry, 2, 231-8 (English) 1989. CODEN: PJPCE9. ISSN: 0915-1699.
- AB O mols. from air plasma were incorporated into

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polypropylene-supported plasma-prepd. hexamethyldisiloxane
     polymers (I), 4-methyl-1-pentene polymers (II),
     octamethylcyclotetrasiloxane polymers (III),
     vinylmethyldiethylsilane polymers (IV), and vinyltrimethylsilane
     polymers (V) creating highly crosslinked polymer membranes
       N-O sepns. by supported plasma-prepd. I, III, IV, and V
    membranes were greater than those by solvent-cast
    membranes. H-N sepns. by supported plasma-prepd.
     II membranes were greater than those by solvent-cast
    membranes at 25-75°. The wettabilities of supported
    plasma-prepd. poly(2-hydroxyethyl methacrylate),
     poly(N-vinylpyrrolidone), and III membranes increased as
     the O content of the membrane surfaces increased with
     increasing polymn. time. The reactivity of carbon fibers towards
     epoxy resin increased as the surface concn. of polar functional
     groups increased due to O-plasma etching.
     1333-74-0, Hydrogen, properties 7782-44-7, Oxygen,
    properties
        (sepn. of, from nitrogen, by polypropylene-supported
        plasma-prepd. polymer membranes)
     1333-74-0 HCA
    Hydrogen (8CI, 9CI) (CA INDEX NAME)
     7782-44-7 HCA
    Oxygen (8CI, 9CI) (CA INDEX NAME)
o = o
    7732-18-5
        (wettability, of plasma-prepd. poly(hydroxyethyl
       methacrylate) and poly(vinylpyrrolidone) and
       poly(octamethylcyclotetrasiloxane) membranes on
       polypropylene, polymn. time and surface oxygen content in
       relation to)
     7732-18-5 HCA
    Water (8CI, 9CI) (CA INDEX NAME)
    36-5 (Physical Properties of Synthetic High Polymers)
    Section cross-reference(s): 35
    plasma polymer membrane gas permeability;
    siloxane plasma membrane gas permeability;
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IT

RN

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H-H

RN

CN

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H<sub>2</sub>O

CC

ST

silane polymer membrane gas permeability; polymethylpentene plasma membrane gas permeability; wettability plasma polymer membrane ; polyhydroxyethyl methacrylate plasma membrane wettability; polyvinylpyrrolidone plasma membrane wettability; carbon fiber reactivity plasma etching IT Plasma, chemical and physical effects (air, polymn. by, of hydrophilic and hydrophobic and organosilicon monomers, oxygen content and wettability in relation to) Wettability IT (of plasma-prepd. poly(hydroxyethyl methacrylate) and poly(vinylpyrrolidone) and poly(octamethylcyclotetrasiloxane) membranes on polypropylene, polymn. time and surface oxygen content in relation to) IT Permeability and Permeation (of polypropylene-supported plasma-prepd. polymer membranes, to hydrogen and nitrogen and oxygen) IT Membranes (polymeric, polypropylene-supported, plasma-prepd., gas permeability and wettability of) Siloxanes and Silicones, uses and miscellaneous IT (polypropylene-supported plasma-prepd. membranes of, gas separability and surface wettability ΙT Epoxy resins, uses and miscellaneous (reactivity of carbon fibers with, effect of oxygenplasma etching of fiber surfaces on) IT Carbon fibers, properties (reactivity of, with epoxy resin, effect of oxygen-plasma etching of fiber surfaces on) IT Functional groups (oxygen-contg., on carbon fibers and polypropylene-supported plasma-prepd. polymer membranes, surface reactivity and wettability in relation to) IT Polymerization (plasma, of hydrophilic and hydrophobic and organosilicon monomers, on polypropylene membranes, oxygen content and wettability in relation to) IT 7440-44-0 (carbon fibers, reactivity of, with epoxy resin, effect of oxygen-plasma etching of fiber surfaces on) 25068-26-2, 4-Methyl-1-pentene homopolymer IT (plasma-prepd. membranes of, on polypropylene, oxygen atoms and crosslinking in, hydrogen and nitrogen gas separability in relation to) IT 26298-61-3, Hexamethyldisiloxane homopolymer 25036-32-2 105218-98-2, Vinylmethyldiethylsilane homopolymer

(plasma-prepd. membranes of, on polypropylene, oxygen atoms and crosslinking in, nitrogen and oxygen gas separability in relation to)

IT 25037-57-4, Octamethylcyclotetrasiloxane homopolymer (plasma-prepd. membranes of, on polypropylene, oxygen atoms and crosslinking in, wettability and nitrogen and oxygen separability in relation to)

IT 9003-39-8, N-Vinylpyrrolidone homopolymer 25249-16-5, 2-Hydroxyethyl methacrylate homopolymer (plasma-prepd. membranes of, on

polypropylene, wettability of)

IT 9003-07-0

(plasma-prepd. polymer membranes supported on, hydrogen and nitrogen and oxygen sepn. by)

IT 128418-38-2, SB 1

(reaction of, with carbon fibers, effect of oxygen-plasma treatment of fiber surfaces on)

IT 1333-74-0, Hydrogen, properties 7782-44-7, Oxygen, properties

(sepn. of, from nitrogen, by polypropylene-supported plasma-prepd. polymer membranes)

IT 7732-18-5

(wettability, of plasma-prepd. poly(hydroxyethyl methacrylate) and poly(vinylpyrrolidone) and poly(octamethylcyclotetrasiloxane) membranes on polypropylene, polymn. time and surface oxygen content in relation to)

- L66 ANSWER 29 OF 32 HCA COPYRIGHT 2006 ACS on STN

  108:122946 Formation of functional tin oxide thin films by
  plasma chemical vapor deposition. Fushimi, Masahiro;
  Shimizu, Isamu (Canon K. K., Japan). Jpn. Kokai Tokkyo Koho JP
  62278268 A2 19871203 Showa, 5 (Japanese). CODEN: JKXXAF.
  APPLICATION: JP 1986-120152 19860527.
- AB The title process is carried out by activation of volatile Sn compds., oxidizing gases, and H(g) by simultaneous irradn. of the gases with the same activation source. The films are useful as transparent electrodes, permselective **membranes** (no data), etc. Thus, a **plasma** was formed using a gas mixt. of SnCl4, O, H, and He to deposit a 1- $\mu$  SnO2 film on a glass substrate. The film showed surface resistivity 10 kΩ and light transmittance >80%.
- IT 7732-18-5, Water, reactions 7782-44-7, Oxygen,
  reactions

(plasma contg., with hydrogen and volatile tin compds.,

transparent conductive tin dioxide film deposition from) 7732-18-5 HCA RNCN Water (8CI, 9CI) (CA INDEX NAME) H<sub>2</sub>O RN 7782-44-7 HCA CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IT 1333-74-0, Hydrogen, reactions (plasma contg., with volatile tin compds. and oxidizing gases, conductive tin dioxide film deposition from) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CNH-HIC ICM C23C016-40 ICS C23C016-50; G02F001-133; H01B013-00; H01L021-28; H01L031-04 76-2 (Electric Phenomena) CC Section cross-reference(s): 42, 49, 75 ST tin oxide conductive transparent film; plasma chem vapor deposition film 7646-78-8, Tin tetrachloride, reactions IT (plasma contg., with hydrogen and oxidizing gases, transparent conductive tin dioxide film deposition from) 7732-18-5, Water, reactions 7782-44-7, Oxygen, IT (plasma contg., with hydrogen and volatile tin compds., transparent conductive tin dioxide film deposition from) IT 1333-74-0, Hydrogen, reactions (plasma contg., with volatile tin compds. and oxidizing gases, conductive tin dioxide film deposition from) 18282-10-5P, Tin dioxide IT (thin films, transparent, elec. conductive, manuf. of, by plasma chem. vapor deposition) ANSWER 30 OF 32 HCA COPYRIGHT 2006 ACS on STN 108:97109 Study of the membrane method of carbon monoxide separation from a carbon dioxide-carbon monoxide-oxygen mixture in a plasmachemical carbonic acid cycle for hydrogen production from water. Vasil'ev, Yu. B.; Kozhevnikov, V. Yu.; Krotov, M. F.; Parovichnikov, A. I.; Sivakov, A. A.; Tyunin, A. N.; Urvachev, N. A.

(USSR). Vopr. Atom. Nauki i Tekhn. Atom.-vodorod. Energ. i Tekhnol., Moskva (2), 50-2 From: Ref. Zh., Khim. 1987, Abstr. No. 21L140 (Russian) 1987. Title only translated.

TT 7782-44-7, Oxygen, uses and miscellaneous
 (carbon monoxide sepn. from carbon dioxide and, in hydrogen
 manuf. from water by plasmachem. cycle)

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME)

o = o

AB

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

IT 1333-74-0P, Hydrogen, preparation (manuf. of, from water by plasmachem. cycle, carbon monoxide membrane sepn. from carbon dioxide and oxygen in)

RN 1333-74-0 HCA

CN Hydrogen (8CI, 9CI) (CA INDEX NAME)

H-H

CC 49-1 (Industrial Inorganic Chemicals)

ST hydrogen manuf water plasmachem cycle; carbon monoxide sepn membrane

IT Plasma, chemical and physical effects
(water decompn. in, hydrogen manuf. by, carbon monoxide sepn.
from carbon dioxide and oxygen in)

TT 7782-44-7, Oxygen, uses and miscellaneous
 (carbon monoxide sepn. from carbon dioxide and, in hydrogen
 manuf. from water by plasmachem. cycle)

IT 1333-74-0P, Hydrogen, preparation (manuf. of, from water by plasmachem. cycle, carbon monoxide membrane sepn. from carbon dioxide and oxygen in)

IT 630-08-0P, Carbon monoxide, preparation (sepn. of, from carbon dioxide and oxygen by membrane

in hydrogen manuf. from water by plasmachem. cycle)

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L66 ANSWER 31 OF 32 HCA COPYRIGHT 2006 ACS on STN
107:29514 Basic processes in glow discharge plasmas.
     Venugopalan, M. (Dep. Chem., West. Illinois Univ., Macomb, IL,
     61455, USA). Nuclear Instruments & Methods in Physics Research,
     Section B: Beam Interactions with Materials and Atoms, B23(4),
     405-17 (English) 1987. CODEN: NIMBEU. ISSN: 0168-583X.
AΒ
     A review with 62 refs. in which the d.c. and radio-
     frequency glow discharge plasmas are
     described in terms of their evolution, mechanism, spatial
     characteristics, and voltage-current relation. The basic
     plasma processes, such as excitation, ionization/dissocn.
     and recombination are reviewed using examples of the chem. reactive
     H2 + O2/H2O and N2 + /NH3 plasmas. This is followed by a discussion
     of some of the heterogeneous processes occurring at the walls and
     substrates, namely film formation, etching, nitriding and analogous
     reactions.
ΙT
     1333-74-0, Hydrogen, properties 7732-18-5,
     Water, vapor 7782-44-7, Oxygen,
     properties
        (plasma contg., basic processes in
        glow discharge plasmas in relation to)
     1333-74-0 HCA
RN
     Hydrogen (8CI, 9CI) (CA INDEX NAME)
CN
H-H
     7732-18-5 HCA
RN
CN
     Water (8CI, 9CI) (CA INDEX NAME)
H20
RN
     7782-44-7 HCA
     Oxygen (8CI, 9CI) (CA INDEX NAME)
CN
0 = 0
CC
     71-0 (Nuclear Technology)
     Section cross-reference(s): 76
     review glow discharge plasma; fusion
ST
    plasma glow discharge review
    Nuclear fusion reactor fuels and plasmas
IT
        (erosion by, basic processes in glow discharge
```

plasmas in relation to)

IT Coating process

(glow discharge plasma basic process in relation to)

IT Plasma

Plasma, chemical and physical effects
 (glow-discharge)

IT Erosion

(of fusion reactor materials, basic processes in **glow discharge** plasmas in relation to)

IT Nuclear fusion reactors

(first walls, erosion of, basic processes in **glow discharge** in relation to)

IT 1333-74-0, Hydrogen, properties 7664-41-7, Ammonia, properties 7727-37-9, Nitrogen, properties 7732-18-5, Water, vapor 7782-44-7, Oxygen, properties

(plasma contg., basic processes in glow discharge plasmas in relation to)

- L66 ANSWER 32 OF 32 HCA COPYRIGHT 2006 ACS on STN
  98:205876 Preliminary design of a fusion reactor fuel cleanup system by the palladium-alloy membrane method. Yoshida, Hiroshi;
  Konishi, Satoshi; Naruse, Yuji (Div. Thermonucl. Fus. Res., Japan At. Energy Res. Inst., Tokai, Japan). Nuclear Technology/Fusion, 3(3), 471-84 (English) 1983. CODEN: NTFUDQ. ISSN: 0272-3921.
- AB A design for a Pd diffuser and fuel cleanup system for a D-T fusion reactor is proposed. The feasibility of the Pd-alloy membrane method is discussed based on earlier studies.

  Operating conditions of the Pd diffuser were detd. exptl.

  Dimensions of the diffuser were estd. from computer simulation. A fuel cleanup system is designed under the feed conditions of the Tritium Systems Test Assembly at Los Alamos National Lab. The system is composed of Pd diffusers, catalytic oxidizer, freezer, and Zn beds and has some advantages in system layout and operation. This design can readily be extended to other conditions of plasma exhaust gases.

TT 7732-18-5, uses and miscellaneous 7782-44-7, uses and miscellaneous

(palladium alloy membrane method for cleanup of fusion reactor fuel system contg.)

RN 7732-18-5 HCA

CN Water (8CI, 9CI) (CA INDEX NAME)

H<sub>2</sub>O

RN 7782-44-7 HCA

CN Oxygen (8CI, 9CI) (CA INDEX NAME) 0 = 0IT1333-74-0, properties (permeability of, in palladium alloy membrane system for fusion reactor fuel cleanup) RN 1333-74-0 HCA Hydrogen (8CI, 9CI) (CA INDEX NAME) CN H-HCC 71-2 (Nuclear Technology) ST fusion reactor fuel cleanup; palladium membrane fuel cleanup IT Membranes (palladium alloy, in nuclear fusion reactor fuel cleanup system) IT Nuclear fusion reactors (fuels and plasmas, palladium alloy membrane method for cleanup of) Palladium alloy, base IT (membrane, in nuclear fusion reactor fuel cleanup 74-82-8, uses and miscellaneous 124-38-9, uses and miscellaneous IT 630-08-0, uses and miscellaneous 7664-41-7, uses and miscellaneous 7727-37-9, uses and miscellaneous **7732-18-5**, uses and miscellaneous 7782-44-7, uses and miscellaneous (palladium alloy membrane method for cleanup of fusion reactor fuel system contg.) ΙT 10028-17-8, uses and miscellaneous (palladium-alloy membrane system for cleanup of fusion reactor fuel contq.) IT **1333-74-0**, properties 7782-39-0, properties (permeability of, in palladium alloy membrane system for fusion reactor fuel cleanup) => D HIS L67-FILE 'HCA' ENTERED AT 14:30:44 ON 20 DEC 2006 25797 S L3 (L) RACT/RL L67 L68 1345 S L13 AND L14 AND L67

2 S L69 NOT (L63 OR L64 OR L65 OR L66)

0 S L70 AND 1840-2003/PRY, PY

L69

L70 L71 25 S L68 AND L25